

(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property
Organization
International Bureau



(43) International Publication Date
10 September 2004 (10.09.2004)

PCT

(10) International Publication Number
WO 2004/076450 A1

(51) International Patent Classification⁷: C07D 471/04, A61K 31/437

(21) International Application Number:
PCT/EP2004/001974

(22) International Filing Date: 27 February 2004 (27.02.2004)

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data:
ES 2003 00640 27 February 2003 (27.02.2003) ES
ES 2003 00727 14 March 2003 (14.03.2003) ES
ES 2003 02504 27 October 2003 (27.10.2003) ES

(71) Applicant (for all designated States except US): J. URI-
ACH Y COMPAÑIA S.A. [ES/ES]; Av. Camí Reial,
51-57, E-08184 Palau-Solità i Plegamans (ES).

(72) Inventors; and

(75) Inventors/Applicants (for US only): ROSALES, Car-
men, Almansa [ES/ES]; C/Teodora Lamadrid, 41,
E-08022 Barcelona (ES). BERNADÓ, Marina, Virgili
[ES/ES]; Plaça Lesseps, 11, E-08023 Barcelona (ES).

(74) Agent: ZUMSTEIN F. and KLINGSEISEN F.;
Bräuhausstr. 4, 80331 München (DE).

(81) Designated States (unless otherwise indicated, for every
kind of national protection available): AF, AG, AI, AM,
AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN,
CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI,
GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE,
KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD,
MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG,
PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM,
TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM,
ZW.

(84) Designated States (unless otherwise indicated, for every
kind of regional protection available): ARIPO (BW, GH,
GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW),
Eurasian (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), Euro-
pean (AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR,
GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK,
TR), OAPI (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW,
ML, MR, NE, SN, TD, TG).

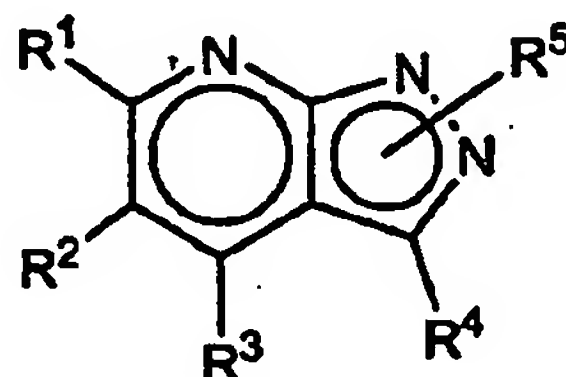
Published:

- with international search report
- before the expiration of the time limit for amending the
claims and to be republished in the event of receipt of
amendments

For two-letter codes and other abbreviations, refer to the "Guid-
ance Notes on Codes and Abbreviations" appearing at the begin-
ning of each regular issue of the PCT Gazette.

WO 2004/076450 A1

(54) Title: PYRAZOLOPYRIDINE DERIVATES



(I)

(57) Abstract: New compounds of formula (I) and the salts, solvates and prodrugs thereof, wherein the meanings for the various
substituents are as disclosed in the description. These compounds are useful as p38 kinase inhibitors.

Pyrazolopyridine derivatives**Field of the invention**

The present invention relates to a new series of pyrazolopyridine derivatives, a process to prepare them, pharmaceutical compositions containing these compounds and their application in medicine.

Background of the invention

Kinases are proteins involved in different cellular responses to external signals. In the Nineties, a new family of kinases called MAPK (*mitogen-activated protein kinases*) was discovered. MAPK activate their substrates by phosphorylation in serine and threonine residues.

MAPK are activated by other kinases in response to a wide range of signals including growth factors, pro-inflammatory cytokines, UV radiation, endotoxins and osmotic stress. Once they are activated, MAPK activate by phosphorylation other kinases or proteins, such as transcription factors, which, ultimately, induce an increase or a decrease in expression of a specific gene or group of genes.

The MAPK family includes kinases such as p38, ERK (*extracellular-regulated protein kinase*) and JNK (*C-Jun N-terminal kinase*).

Kinase p38 plays a crucial role in cellular response to stress and in the activation pathway in the synthesis of numerous cytokines, especially tumor necrosis factor (TNF- α), interleukin-1 (IL-1), interleukin-6 (IL-6) and interleukin-8 (IL-8).

IL-1 and TNF- α are produced by macrophages and monocytes and are involved in the mediation of immunoregulation processes and other physiopathological conditions. For example, elevated levels of TNF- α are associated with inflammatory and autoimmune diseases and with processes that trigger the degradation of connective and bone tissue such as rheumatoid arthritis, osteoarthritis, diabetes, inflammatory bowel disease and sepsis.

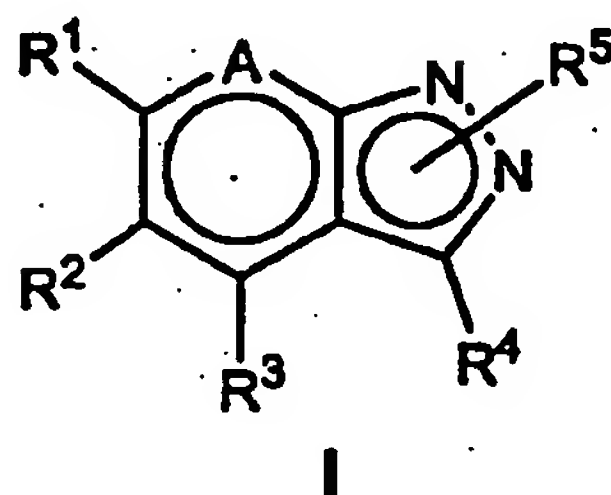
Thus, p38 kinase inhibitors could be useful to treat or prevent diseases mediated by cytokines such as IL-1 and TNF- α , as mentioned previously.

On the other hand, it has also been found that p38 inhibitors inhibit other pro-inflammatory proteins such as IL-6, IL-8, interferon- γ and GM-CSF (*granulocyte-macrophage colony-stimulating factor*). Moreover, in recent studies it has been found that p38 inhibitors not only block cytokine synthesis but also the

cascade of signals that these induce, such as induction of the cyclooxygenase-2 enzyme (COX-2).

Description of the Invention

One aspect of the present invention relates to the new compounds of
5 general formula I



wherein:

A represents N or N⁺O⁻;

10 R¹ represents phenyl or Het optionally substituted with one or more substituents selected from R^a and R^b;

R² represents Het optionally substituted with one or more substituents selected from R^a and R^b;

15

R³ represents H, Cy optionally substituted with one or more substituents selected from R^a and R^b, or R³ represents C₁₋₆alkyl optionally substituted with one or more substituents selected from R^b and Cy*, wherein Cy* can be optionally substituted with one or more substituents selected from R^b and R^c;

20

R⁴ represents H, R^a, halogen, -OR^a, -OCOR^a, -OSO₂R^a, -OCONR^aR^a, -NO₂, -CN, -COR^a, -CO₂R^a, -CONR^aR^a, -NR^aR^a, -NR^aCOR^a, -NR^aCONR^aR^a, -NR^aCO₂R^a, -NR^aSO₂R^a, -SR^a, -SOR^a, -SO₂R^a or -SO₂NR^aR^a;

25 R⁵ can be placed on any of the 2 N of the pyrazole ring of formula I and represents H or R^f;

each R^a independently represents C_{1-6} alkyl, C_{2-6} alkenyl, C_{2-6} alkynyl or Cy, wherein the groups C_{1-6} alkyl, C_{2-6} alkenyl or C_{2-6} alkynyl can be optionally substituted with one or more substituents selected from R^b and Cy^* , and wherein any of the groups Cy or Cy^* can be optionally substituted with one or more substituents selected
5 from R^b and R^c ;

each R^a independently represents H or R^a ;

each R^b independently represents halogen, $-OR^c$, $-OCOR^c$, $-OSO_2R^c$,
10 $-OCONR^cR^c$, $-NO_2$, $-CN$, $-COR^c$, $-CO_2R^c$, $-CONR^cR^c$, $-CONR^cNR^cR^c$,
 $-NR^cR^c$, $-NR^cCOR^c$, $-NR^cCONR^cR^c$, $-NR^cCO_2R^c$, $-NR^cSO_2R^c$, $-SR^c$, $-SOR^c$,
 $-SO_2R^c$, $-SO_2NR^cR^c$, $-C(NR^c)NR^cR^c$, $-C(NSO_2NR^cR^c)NR^cR^c$, $-C(NOR^c)R^c$,
 $-C(NNR^cR^c)R^c$, $-NR^cC(NR^c)NR^cR^c$ or $-NR^cC(NCN)NR^cR^c$;

15 each R^c independently represents C_{1-6} alkyl, C_{2-6} alkenyl, C_{2-6} alkynyl or Cy, wherein all these groups can be optionally substituted with one or more substituents R^d ;

each R^c independently represents H or R^c ;

20 each R^d independently represents halogen, R^e , $-OR^e$, $-OCOR^e$, $-OSO_2R^e$,
 $-OCONR^eR^e$, $-NO_2$, $-CN$, $-COR^e$, $-CO_2R^e$, $-CONR^eR^e$, $-CONR^eNR^eR^e$,
 $-NR^eR^e$, $-NR^eCOR^e$, $-NR^eCONR^eR^e$, $-NR^eCO_2R^e$, $-NR^eSO_2R^e$, $-SR^e$, $-SOR^e$,
 $-SO_2R^e$, $-SO_2NR^eR^e$, $-C(NR^e)NR^eR^e$, $-C(NSO_2NR^eR^e)NR^eR^e$, $-C(NOR^e)R^e$,
 $-C(NNR^eR^e)R^e$, $-NR^eC(NR^e)NR^eR^e$, $-NR^eC(NCN)NR^eR^e$ or Cy optionally
25 substituted with one or more substituents selected from halogen, R^e , $-OR^e$,
 $-OCOR^e$, $-OSO_2R^e$, $-OCONR^eR^e$, $-NO_2$, $-CN$, $-COR^e$, $-CO_2R^e$, $-CONR^eR^e$,
 $-CONR^eNR^eR^e$, $-NR^eR^e$, $-NR^eCOR^e$, $-NR^eCONR^eR^e$, $-NR^eCO_2R^e$,
 $-NR^eSO_2R^e$, $-SR^e$, $-SOR^e$, $-SO_2R^e$, $-SO_2NR^eR^e$, $-C(NR^e)NR^eR^e$,
 $-C(NSO_2NR^eR^e)NR^eR^e$, $-C(NOR^e)R^e$, $-C(NNR^eR^e)R^e$, $-NR^eC(NR^e)NR^eR^e$ and
30 $-NR^eC(NCN)NR^eR^e$;

each R^e independently represents C_{1-6} alkyl or halo C_{1-6} alkyl;

each R^e independently represents H or R^e ;

R^f represents C_{1-6} alkyl, C_{2-6} alkenyl, C_{2-6} alkynyl or Cy, wherein the groups C_{1-6} alkyl, C_{2-6} alkenyl or C_{2-6} alkynyl can be optionally substituted with one or more
 5 substituents selected from R^g and Cy^* , and wherein any of the groups Cy or Cy^* can be optionally substituted with one or more substituents selected from R^g and R^h ;

each R^g independently represents halogen, $-OR^a$, $-OCOR^a$, $-OSO_2R^a$,
 10 $-OCONR^aR^a$, $-NO_2$, $-CN$, $-COR^a$, $-CO_2R^a$, $-CONR^aR^a$, $-CONR^aNR^aR^a$,
 $-NR^aR^a$, $-NR^aCOR^a$, $-NR^aCONR^aR^a$, $-NR^aCO_2R^a$, $-NR^aSO_2R^a$, $-SR^a$, $-SOR^a$,
 $-SO_2R^a$, $-SO_2NR^aR^a$, $-C(NR^a)NR^aR^a$, $-C(NSO_2NR^aR^a)NR^aR^a$, $-C(NOR^a)R^a$,
 $-C(NNR^aR^a)R^a$, $-NR^aC(NR^a)NR^aR^a$ or $-NR^aC(NCN)NR^aR^a$;

15 Het in the above definitions represents pyridine, pyrazine, pyrimidine, pyridazine,
 2(1H)-pyridone, 2(1H)-pyrazinone, 2(1H)-pyrimidinone or 2(1H)-pyridazinone;

Cy or Cy^* in the above definitions represent a partially unsaturated, saturated or
 aromatic 3- to 7-membered monocyclic or 8- to 12-membered bicyclic carbocyclic
 20 ring, which optionally contains from 1 to 4 heteroatoms selected from N, S and O,
 which can optionally contain 1 or 2 oxo groups when the ring is saturated or
 partially unsaturated, and wherein said ring or rings can be bonded to the rest of
 the molecule through a carbon or a nitrogen atom.

The present invention also relates to the addition salts of the compounds of
 25 the invention as well as their solvates and prodrugs. A prodrug is defined as any
 precursor of a compound of formula I that can be transformed in vivo into a
 compound of formula I.

Some compounds of formula I can have chiral centres that can give rise to
 various stereoisomers. The present invention relates to each of these
 30 stereoisomers and also mixtures thereof. Moreover, some of the compounds of
 the present invention can show cis/trans isomers. The present invention relates to
 each of the geometric isomers and mixtures thereof.

The compounds of formula I are selective p38 kinase inhibitors.

Thus, another aspect of this invention relates to the pharmaceutical compositions which comprise an effective amount of a compound of formula I or a pharmaceutically acceptable salt, solvate or prodrug thereof and one or more pharmaceutically acceptable excipients.

5 Another aspect of the present invention relates to the use of a compound of formula I or a pharmaceutically acceptable salt, solvate or prodrug thereof for the manufacture of a medicament for the treatment or prevention of diseases mediated by p38.

10 Another aspect of the present invention relates to the use of a compound of formula I or a pharmaceutically acceptable salt, solvate or prodrug thereof for the manufacture of a medicament for the treatment or prevention of diseases mediated by cytokines.

15 Another aspect of the present invention relates to the use of a compound of formula I or a pharmaceutically acceptable salt, solvate or prodrug thereof for the manufacture of a medicament for the treatment or prevention of diseases mediated by TNF- α , IL-1, IL-6 and/or IL-8.

20 Another aspect of the present invention relates to the use of a compound of formula I or a pharmaceutically acceptable salt, solvate or prodrug thereof for the manufacture of a medicament for the treatment or prevention of a disease selected from immune, autoimmune and inflammatory diseases, cardiovascular diseases, infectious diseases, bone resorption disorders, neurodegenerative diseases, proliferative diseases and processes associated with the induction of cyclooxygenase-2.

25 Another aspect of the present invention relates to the use of a compound of formula I or a pharmaceutically acceptable salt, solvate or prodrug thereof for the treatment or prevention of diseases mediated by p38.

Another aspect of the present invention relates to the use of a compound of formula I or a pharmaceutically acceptable salt, solvate or prodrug thereof for the treatment or prevention of diseases mediated by cytokines.

30 Another aspect of the present invention relates to the use of a compound of formula I or a pharmaceutically acceptable salt, solvate or prodrug thereof for the treatment or prevention of diseases mediated by TNF- α , IL-1, IL-6 and/or IL-8.

Another aspect of the present invention relates to the use of a compound of

formula I or a pharmaceutically acceptable salt, solvate or prodrug thereof for the treatment or prevention of a disease selected from immune, autoimmune and inflammatory diseases, cardiovascular diseases, infectious diseases, bone resorption disorders, neurodegenerative diseases, proliferative diseases and processes associated with the induction of cyclooxygenase-2.

Another aspect of the present invention relates to a method of treating or preventing diseases mediated by p38 in a subject in need thereof, especially a human being, which comprises administering to said subject a therapeutically effective amount of a compound of formula I or a pharmaceutically acceptable salt, solvate or prodrug thereof.

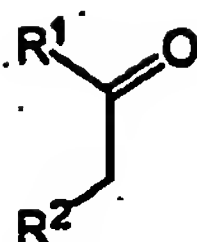
Another aspect of the present invention relates to a method of treating or preventing diseases mediated by cytokines in a subject in need thereof, especially a human being, which comprises administering to said subject a therapeutically effective amount of a compound of formula I or a pharmaceutically acceptable salt, solvate or prodrug thereof.

Another aspect of the present invention relates to a method of treating or preventing diseases mediated by TNF- α , IL-1, IL-6 and/or IL-8 in a subject in need thereof, especially a human being, which comprises administering to said subject a therapeutically effective amount of a compound of formula I or a pharmaceutically acceptable salt, solvate or prodrug thereof.

Another aspect of the present invention relates to a method of treating or preventing a disease selected from immune, autoimmune and inflammatory diseases, cardiovascular diseases, infectious diseases, bone resorption disorders, neurodegenerative diseases, proliferative diseases and processes associated with the induction of cyclooxygenase-2 in a subject in need thereof, especially a human being, which comprises administering to said subject a therapeutically effective amount of a compound of formula I or a pharmaceutically acceptable salt, solvate or prodrug thereof.

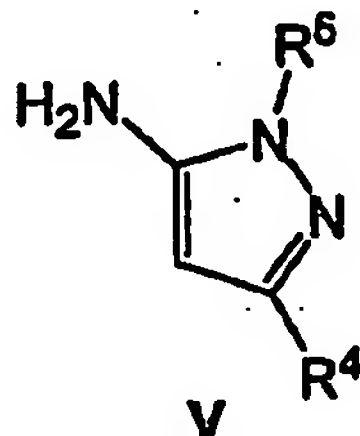
Another aspect of the present invention relates to a process for the preparation of a compound of formula I, which comprises:
(a) reacting a ketone of formula IV

7



IV

wherein R^1 and R^2 have the meaning described above, with an aminopyrazole of formula V and an aldehyde of formula VI

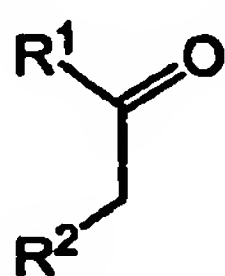


V

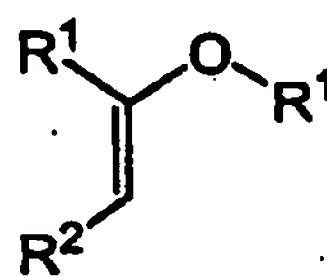


VI

- 5 wherein R^3 , R^4 and R^5 have the meaning described above; or
 (b) when in a compound of formula I, R^5 represents H and R^3 has the same meaning as R^1 , reacting a ketone of formula IV or an enolate of formula VII

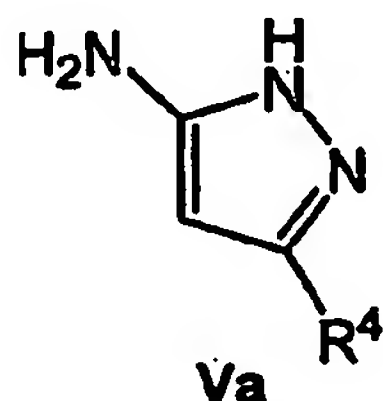


IV



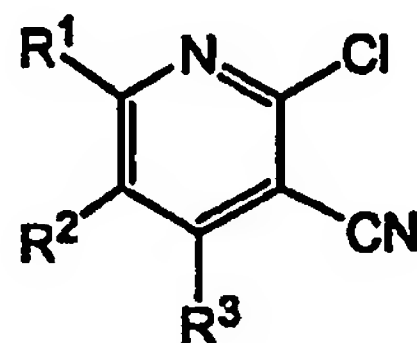
VII

- 10 wherein R^1 and R^2 have the meaning described above, with an aminopyrazole of formula Va



Va

wherein R^4 has the meaning described above; or
 (c) when in a compound of formula I R^4 represents NH_2 , treating a compound of formula XIX

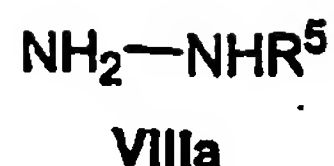


XIX

15

wherein R^1 , R^2 and R^3 have the meaning described above, with a hydrazine of

formula VIIIa



wherein R^5 has the meaning described above; or

- (d) converting, in one or a plurality of steps, a compound of formula I into another compound of formula I; and
- (e) if desired, after the previous steps, reacting a compound of formula I with a base or an acid to give the corresponding salt.

In the previous definitions, the term C_{1-6} alkyl, as a group or part of a group, means a straight or branched alkyl chain which contains from 1 to 6 carbon atoms. Examples include among others the groups methyl, ethyl, propyl, isopropyl, butyl, isobutyl, *sec*-butyl, *tert*-butyl, pentyl, isopentyl, neopentyl and hexyl.

A halo C_{1-6} alkyl group means a group resulting from the replacement of one or more hydrogen atoms from a C_{1-6} alkyl group with one or more halogen atoms (i.e. fluoro, chloro, bromo or iodo), which can be the same or different. Examples include, among others, trifluoromethyl, fluoromethyl, 1-chloroethyl, 2-chloroethyl, 1-fluoroethyl, 2-fluoroethyl, 2-bromoethyl, 2-iodoethyl, 2,2,2-trifluoroethyl, pentafluoroethyl, 3-fluoropropyl, 3-chloropropyl, 2,2,3,3-tetrafluoropropyl, 2,2,3,3,3-pentafluoropropyl, heptafluoropropyl, 4-fluorobutyl, nonafluorobutyl, 5-fluoropentyl and 6-fluorohexyl.

The term C_{2-6} alkenyl, as a group or part of a group, means a straight or branched alkyl chain which contains from 2 to 6 carbon atoms and that also contains one or more double bonds. Examples include, among others, the groups ethenyl, 1-propenyl, 2-propenyl, isopropenyl, 1-butenyl, 2-butenyl, 3-butenyl, 1,3-butadienyl, 1-pentenyl, 2-pentenyl, 3-pentenyl, 4-pentenyl, 1-hexenyl, 2-hexenyl, 3-hexenyl, 4-hexenyl and 5-hexenyl.

The term C_{2-6} alkynyl, as a group or part of a group, means a straight or branched alkyl chain which contains from 2 to 6 carbon atoms and that also contains one or more triple bonds. Examples include the groups ethynyl, 1-propynyl, 2-propynyl, 1-butylnyl, 2-butylnyl, 3-butylnyl, 1,3-butadiynyl, 1-pentylnyl, 2-pentylnyl, 3-pentylnyl, 4-pentylnyl, 1-hexynyl, 2-hexynyl, 3-hexynyl, 4-hexynyl and 5-hexynyl.

An oxo group means a carbonyl group ($-\text{CO}-$).

A halogen radical means fluoro, chloro, bromo or iodo.

Het in the definitions of R^1 and R^2 means pyridine, pyrazine, pyrimidine, pyridazine, 2(1H)-pyridone, 2(1H)-pyrazinone, 2(1H)-pyrimidinone or 2(1H)-pyridazinone. As mentioned previously, these groups can be optionally substituted
5 with one or more substituents selected from R^a and R^b , which can be placed on any available position of the Het group, and can be bonded to the rest of the molecule via any available carbon or nitrogen atom.

The term Cy or Cy^* , as a group or part of a group, means a 3- to 7-membered monocyclic carbocyclic group or an 8- to 12-membered bicyclic
10 carbocyclic group which can be partially unsaturated, saturated or aromatic and which can optionally contain from 1 to 4 heteratoms selected from N, S and O. When the Cy or Cy^* group is saturated or partially unsaturated, it can optionally contain 1 or 2 oxo groups. The Cy or Cy^* ring or rings can be substituted as mentioned in the definition of general formula I, these substituents being placed on
15 any available position, and can be bonded to the rest of the molecule through any available carbon or nitrogen atom. Examples of Cy or Cy^* groups include, among others, cyclopropane, cyclobutane, cyclopentane, cyclohexane, cycloheptane, aziridine, oxirane, oxetane, imidazolidine, isothiazolidine, isoxazolidine, oxazolidine, pyrazolidine, pyrrolidine, thiazolidine, dioxane, morpholine,
20 piperazine, piperidine, pyran, tetrahydropyran, azepine, oxazine, oxazoline, pyrrolidine, thiazoline, pyrazoline, imidazoline, isoxazoline, isothiazoline, phenyl, naphthyl, 1,2,4-oxadiazole, 1,2,4-thiadiazole, 1,3,4-oxadiazole, 1,3,4-thiadiazole, furan, imidazole, isoxazole, isothiazole, oxazole, pyrazole, pyrrole, thiazole, thiophene, 1,2,3-triazole, 1,2,4-triazole, pyrazine, pyridazine, pyridine, pyrimidine,
25 benzimidazole, benzofuran, benzothiazole, benzothiofene, imidazopyrazine, imidazopyridazine, imidazopyridine, imidazopyrimidine, indazole, indole, isoindole, isoquinoline, tetrahydroisoquinoline, naphthyridine, pyrazolopyrazine, pyrazolopyridine, pyrazolopyrimidine, purine, quinazoline, quinoline, quinoxaline, cyclobutanone, cyclopentanone, cyclohexanone, cycloheptanone, pyrrolidin-2-one, piperidin-2-one, piperidin-4-one, 2(1H)-pyridone, 2(1H)-pyrazinone, 2(1H)-pyrimidinone, 2(1H)-pyridazinone and phthalimide.
30

The term heteroaryl, as a group or part of a group, means an aromatic 5- or 6-membered monocyclic or 8- to 12-membered bicyclic ring which contains from 1

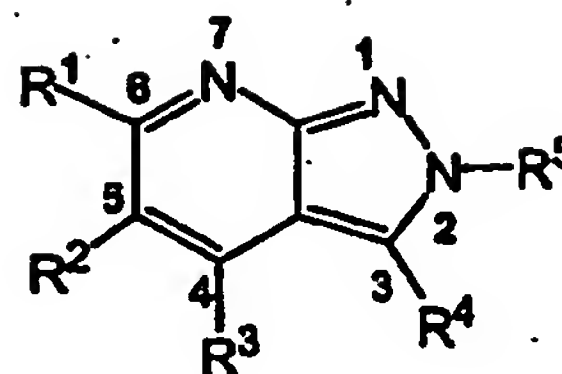
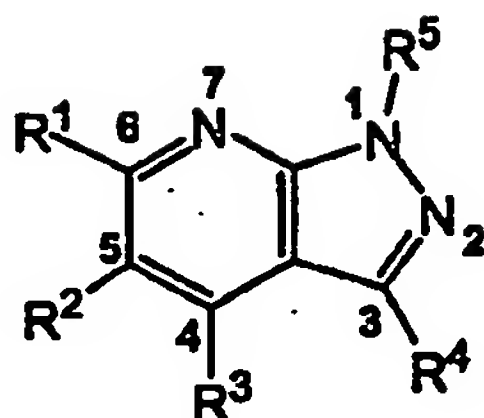
to 4 heteroatoms selected from N, S and O and which can be optionally substituted as disclosed whenever this term is used, wherein said substituents can be placed on any available position. The heteroaryl group can be bonded to the rest of the molecule through any available carbon or nitrogen atom. Examples of heteroaryl groups include among others 1,2,4-oxadiazole, 1,2,4-thiadiazole, 1,3,4-oxadiazole, 1,3,4-thiadiazole, furan, imidazole, isoxazole, isothiazole, oxazole, pyrazole, pyrrole, thiazole, thiophene, 1,2,3-triazole, 1,2,4-triazole, pyrazine, pyridazine, pyridine, pyrimidine, benzimidazole, benzofuran, benzothiazole, benzothiophene, imidazopyrazine, imidazopyridazine, imidazopyridine, imidazopyrimidine, indazole, indole, isoindole, isoquinoline, naphthiridine, pyrazolopyrazine, pyrazolopyridine, pyrazolopyrimidine, purine, quinazoline, quinoline and quinoxaline.

In the previous definitions of Het, heteroaryl, Cy and Cy*, the term is meant to be the radical derived from the corresponding cycle.

In the previous definitions of heteroaryl, Cy and Cy*, when the specified examples refer to a bicycle in general terms, all possible dispositions of the atoms are included. For example, the term pyrazolopyridine is to be understood as including groups such as 1H-pyrazolo[3,4-b]pyridine, pyrazolo[1,5-a]pyridine, 1H-pyrazolo[3,4-c]pyridine, 1H-pyrazolo[4,3-c]pyridine and 1H-pyrazolo[4,3-b]pyridine; the term imidazopyrazine is to be understood as including groups such as 1H-imidazo[4,5-b]pyrazine, imidazo[1,2-a]pyrazine and imidazo[1,5-a]pyrazine and the term pyrazolopyrimidine is to be understood as including groups such as 1H-pyrazolo[3,4-d]pyrimidine, 1H-pyrazolo[4,3-d]pyrimidine, pyrazolo[1,5-a]pyrimidine and pyrazolo[1,5-c]pyrimidine.

The expression "optionally substituted with one or more" means that a group can be substituted with one or more, preferably with 1, 2, 3 or 4 substituents, provided that this group has 1, 2, 3 or 4 positions susceptible of being substituted.

In the previous definitions, when it is mentioned that R⁵ can be placed on any one of the 2N of the pyrazole ring this means that R⁵ can be placed on the N at position 1 of the ring or on the N at position 2. Thus, the compounds of formula I include the following two types of compounds:



An embodiment of the invention are those compounds of formula I as defined above wherein A represents N.

- 5 Another embodiment of the invention are the compounds of formula I wherein A represents N and R⁵ can be placed on any of the 2 N of the pyrazole ring of formula I and represents H or R^a.

- Another embodiment of the invention are the compounds of formula I wherein when R³ and R⁵ both represent H and R² represents Het optionally substituted with one or more substituents selected from halogen, -CN, -CF₃, -OH, -NO₂, -OR^b, -NR^bR^b, -OCF₃, C₁₋₆alkyl, C₂₋₆alkenyl, C₂₋₆alkynyl and Cy, wherein Cy can be optionally substituted with one or more substituents selected from R^b and R^c, and wherein R^b represents C₁₋₆alkyl, then R⁴ is not -NR^aCOR^a, -NHCONHR^a or -NHCO₂R^a.

- 15 Another embodiment of the invention are the compounds of formula I wherein when R³ and R⁵ both represent H, then R⁴ is not -NR^aCOR^a, -NHCONHR^a or -NHCO₂R^a.

- Another embodiment of the invention are the compounds of formula I wherein A represents N; R⁴ represents H, R^a, halogen, -OR^a, -OCOR^a, -OSO₂R^a, -OCONR^aR^a, -NO₂, -CN, -COR^a, -CO₂R^a, -CONR^aR^a, -NR^aR^a, -NR^aSO₂R^a, -SR^a, -SOR^a, -SO₂R^a or -SO₂NR^aR^a; and R⁵ can be placed on any of the 2 N of the pyrazole ring of formula I and represents H or R^a.

- Another embodiment of the invention are the compounds of formula I wherein R¹ represents pyridine or phenyl, wherein all these groups can be optionally substituted with one or more substituents selected from R^a and R^b.

Another embodiment of the invention are the compounds of formula I wherein R¹ represents phenyl optionally substituted with one or more substituents selected from R^a and R^b.

Another embodiment of the invention are the compounds of formula I wherein R^1 represents phenyl optionally substituted with one or more substituents selected from halogen, $-OR^c$, $-NO_2$, $-CN$, $-CONR^cR^c$, $-NR^cR^c$ and C_{1-6} alkyl optionally substituted with one or more substituents selected from halogen, $-OR^c$, $-COR^c$, $-NR^cR^c$ and $-NR^cCOR^c$.

Another embodiment of the invention are the compounds of formula I wherein R^1 represents phenyl optionally substituted with one or more substituents selected from halogen and halo C_{1-6} alkyl.

Another embodiment of the invention are the compounds of formula I wherein R^2 represents pyridine or pyrimidine, wherein all these groups can be optionally substituted with one or more substituents selected from R^a and R^b .

Another embodiment of the invention are the compounds of formula I wherein R^2 represents 4-pyridine or 4-pyrimidine, wherein all these groups can be optionally substituted with one or more substituents selected from R^a and R^b .

Another embodiment of the invention are the compounds of formula I wherein R^2 represents 4-pyridine or 4-pyrimidine, wherein all these groups can be optionally substituted with one or more substituents selected from halogen, $-OR^c$, $-NR^cR^c$, $-SR^c$ and $-SO_2R^c$.

Another embodiment of the invention are the compounds of formula I wherein R^2 represents 4-pyridine.

Another embodiment of the invention are the compounds of formula I wherein R^2 represents 4-pyrimidine substituted with $-NR^cR^c$, wherein in R^2 :
each R^c independently represents H or R^c ;
each R^c independently represents C_{1-6} alkyl optionally substituted with one or more substituents selected from Cy and $-OR^e$; and
each R^e independently represents H or R^e .

Another embodiment of the invention are the compounds of formula I wherein R^3 represents H or Cy optionally substituted with one or more substituents selected from R^a and R^b .

Another embodiment of the invention are the compounds of formula I wherein R^3 represents H, heteroaryl or phenyl, wherein all these groups can be optionally substituted with one or more substituents selected from R^a and R^b .

Another embodiment of the invention are the compounds of formula I wherein R^3 represents heteroaryl or phenyl, wherein all these groups can be optionally substituted with one or more substituents selected from R^a and R^b .

Another embodiment of the invention are the compounds of formula I
5 wherein R^3 represents monocyclic heteroaryl or phenyl, wherein all these groups can be optionally substituted with one or more substituents selected from halogen, $-NO_2$, $-OR^c$, C_{1-6} alkyl and Cy, wherein C_{1-6} alkyl can be optionally substituted with one or more substituents selected from R^b and Cy^* , and any of the groups Cy or Cy^* can be optionally substituted with one or more substituents selected from R^b
10 and R^c .

Another embodiment of the invention are the compounds of formula I wherein R^3 represents monocyclic heteroaryl or phenyl, wherein all these groups can be optionally substituted with one or more substituents selected from halogen, $-NO_2$, $-OR^c$, C_{1-6} alkyl, halo C_{1-6} alkyl and Cy; and wherein in R^3 :
15 each R^c independently represents H or R^c ;
each R^c independently represents C_{1-6} alkyl optionally substituted with one or more substituents R^d ; and
each R^d independently represents Cy.

Another embodiment of the invention are the compounds of formula I
20 wherein R^4 represents H, R^a , halogen, $-OR^a$, $-CN$, $-CONR^aR^a$, $-NR^aR^a$ or $-NR^aCOR^a$.

Another embodiment of the invention are the compounds of formula I wherein R^4 represents H.

Another embodiment of the invention are the compounds of formula I
25 wherein R^5 represents H or R^5 represents R^f and is placed on the N at position 2 of the pyrazole ring.

Another embodiment of the invention are the compounds of formula I wherein R^5 represents R^f and is placed on the N at position 2 of the pyrazole ring.

Another embodiment of the invention are the compounds of formula I
30 wherein R^f represents C_{1-6} alkyl, C_{2-6} alkenyl or Cy, wherein the groups C_{1-6} alkyl or C_{2-6} alkenyl can be optionally substituted with one or more substituents selected from R^g and Cy^* , and wherein any of the groups Cy or Cy^* can be optionally substituted with one or more substituents selected from R^g and R^a .

Another embodiment of the invention are the compounds of formula I wherein R^f represents C_{1-6} alkyl, C_{2-6} alkenyl or Cy, wherein the groups C_{1-6} alkyl or C_{2-6} alkenyl can be optionally substituted with one or more substituents selected from R^g and Cy^* , and wherein any of the groups Cy or Cy^* can be optionally substituted with one or more substituents selected from R^g and R^a , wherein each R^g independently represents halogen, $-OR^a$, $-COR^a$, $-CO_2R^a$, $-CONR^aR^a$, $-NR^aR^a$, $-NR^aCOR^a$, $-NR^aCONR^aR^a$, $-NR^aSO_2R^a$, $-SR^a$, $-SOR^a$ or $-SO_2R^a$.

Another embodiment of the invention are the compounds of formula I wherein R^f represents C_{1-6} alkyl, C_{2-6} alkenyl or Cy, wherein the groups C_{1-6} alkyl or C_{2-6} alkenyl can be optionally substituted with one or more substituents selected from R^g and Cy^* , and wherein any of the groups Cy or Cy^* can be optionally substituted with one or more substituents selected from R^g and R^a , wherein in R^f :

each R^g independently represents $-OR^a$, $-COR^a$, $-CONR^aR^a$, $-NR^aR^a$, $-NR^aCOR^a$, $-NR^aCONR^aR^a$, $-NR^aSO_2R^a$, $-SOR^a$ or $-SO_2R^a$;

each R^a independently represents H or R^a ;

each R^a independently represents Cy or C_{1-6} alkyl, wherein C_{1-6} alkyl can be optionally substituted with one or more substituents selected from R^b and Cy^* , and any of the groups Cy or Cy^* can be optionally substituted with one or more substituents selected from R^b and R^c ;

each R^b independently represents $-OR^c$, $-NR^cR^c$, $-CN$, $-COR^c$, $-SR^c$ or $-SOR^c$;

each R^c independently represents H or R^c ;

each R^b independently represents C_{1-6} alkyl or Cy, wherein all these groups can be optionally substituted with one or more R^d .

Another embodiment of the invention are the compounds of formula I wherein R^5 is placed on the N at position 2 of the pyrazole ring and represents R^f , wherein R^f represents C_{1-6} alkyl optionally substituted with one or more substituents selected from $-OR^a$, $-COR^a$, $-CONR^aR^a$, $-NR^aR^a$, $-NR^aCOR^a$, $-NR^aCONR^aR^a$, $-NR^aSO_2R^a$ and Cy^* optionally substituted with one or more substituents selected from R^a ; wherein in R^f :

each R^a independently represents H or R^a ;

each R^a independently represents Cy or C_{1-6} alkyl, wherein C_{1-6} alkyl can be optionally substituted with one or more substituents selected from R^b and Cy^* , and

any of the groups Cy or Cy* can be optionally substituted with one or more substituents selected from R^b and R^c;

each R^b independently represents -OR^c, -NR^cR^c, -CN, -COR^c, -SR^c or -SOR^c;

each R^c independently represents H or R^c;

5 each R^c independently represents C₁₋₆alkyl or Cy, wherein all these groups can be optionally substituted with one or more R^d; and

each R^d independently represents -OR^e.

Another embodiment of the invention are the compounds of formula I wherein R⁵ represents H or R⁵ represents R^a and is placed on the N at position 2
10 of the pyrazole ring.

Another embodiment of the invention are the compounds of formula I wherein R⁵ represents R^a and is placed on the N at position 2 of the pyrazole ring.

Another embodiment of the invention are the compounds of formula I wherein R^a in R⁵ represents C₁₋₆alkyl, C₂₋₆alkenyl or Cy, wherein the groups
15 C₁₋₆alkyl or C₂₋₆alkenyl can be optionally substituted with one or more substituents selected from R^b and Cy*, and wherein any of the groups Cy or Cy* can be optionally substituted with one or more substituents selected from R^b and R^c.

Another embodiment of the invention are the compounds of formula I wherein R^a in R⁵ represents C₁₋₆alkyl, C₂₋₆alkenyl or Cy, wherein the groups
20 C₁₋₆alkyl or C₂₋₆alkenyl can be optionally substituted with one or more substituents selected from R^b and Cy*, and wherein any of the groups Cy or Cy* can be optionally substituted with one or more substituents selected from R^b and R^c, wherein each R^b in R⁵ independently represents halogen, -OR^c, -COR^c, -CO₂R^c,
-CONR^cR^c, -NR^cR^c, -NR^cCOR^c, -NR^cCONR^cR^c, -NR^cSO₂R^c, -SR^c, -SOR^c or
25 -SO₂R^c.

Another embodiment of the invention are the compounds of formula I wherein R^a in R⁵ represents C₁₋₆alkyl, C₂₋₆alkenyl or Cy, wherein the groups
C₁₋₆alkyl or C₂₋₆alkenyl can be optionally substituted with one or more substituents selected from R^b and Cy*, and wherein any of the groups Cy or Cy* can be
30 optionally substituted with one or more substituents selected from R^b and R^c, wherein in R⁵:

each R^b independently represents -OR^c, -COR^c, -CONR^cR^c, -NR^cR^c, -NR^cCOR^c,
-NR^cCONR^cR^c, -NR^cSO₂R^c, -SOR^c or -SO₂R^c;

each R^c independently represents H or R^c ;

each R^c independently represents Cy or C_{1-6} alkyl, wherein all these groups can be optionally substituted with one or more substituents selected from R^d ;

each R^d independently represents R^e , $-OR^e$, $-NR^eR^e$, $-CN$, $-COR^e$, $-SR^e$, $-SOR^e$

5. or Cy.

Another embodiment of the invention are the compounds of formula I wherein R^5 is placed on the N at position 2 of the pyrazole ring and represents R^a , wherein R^a in R^5 represents C_{1-6} alkyl optionally substituted with one or more substituents selected from $-OR^c$, $-COR^c$, $-CONR^cR^c$, $-NR^cR^c$, $-NR^cCOR^c$,
10 $-NR^cCONR^cR^c$, $-NR^cSO_2R^c$ and Cy* optionally substituted with one or more substituents selected from R^c ; wherein in R^5 :

each R^c independently represents H or R^c ;

each R^c independently represents Cy or C_{1-6} alkyl, wherein all these groups can be optionally substituted with one or more substituents selected from R^d ;

15 each R^d independently represents $-OR^e$, $-NR^eR^e$, $-CN$, $-COR^e$, $-SR^e$, $-SOR^e$ or Cy;

each R^e independently represents H or R^e ; and

each R^e independently represents C_{1-6} alkyl.

Furthermore, all possible combinations of the above-mentioned
20 embodiments form also part of this invention.

The compounds of the present invention may contain one or more basic nitrogens and may, therefore, form salts with organic or inorganic acids that also form part of this invention. Examples of these salts include: salts with inorganic acids such as hydrochloric acid, hydrobromic acid, hydroiodic acid, nitric acid,
25 perchloric acid, sulfuric acid or phosphoric acid, and salts with organic acids such as methanesulfonic acid, trifluoromethanesulfonic acid, ethanesulfonic acid, benzenesulfonic acid, *p*-toluenesulfonic acid, fumaric acid, oxalic acid, acetic acid or malic acid, among others. The compounds of the present invention may contain one or more acidic protons and, therefore, they may also form salts with bases,
30 which also form part of the present invention. Examples of these salts include: salts with inorganic cations such as sodium, potassium, calcium, magnesium, lithium, aluminium, zinc, etc; and salts formed with pharmaceutically acceptable amines such as ammonia, alkylamines, hydroxyalkylamines, lysine, arginine, *N*-

methyglucamine, procaine and the like b. There is no limitation on the type of salt that can be used provided that these are pharmaceutically acceptable when they are used for therapeutic purposes. Salts can be prepared by treating the compound of formula I with a sufficient amount of the desired acid or base to give
5 a salt in the conventional manner. The compounds of formula I and their salts differ in some physical properties but they are equivalent for the purposes of the present invention.

Some of the compounds of the present invention can exist in solvated form, including hydrated forms. In general, the solvated forms with pharmaceutically
10 acceptable solvents such as water, ethanol and the like are equivalent to the unsolvated form for the purposes of the invention.

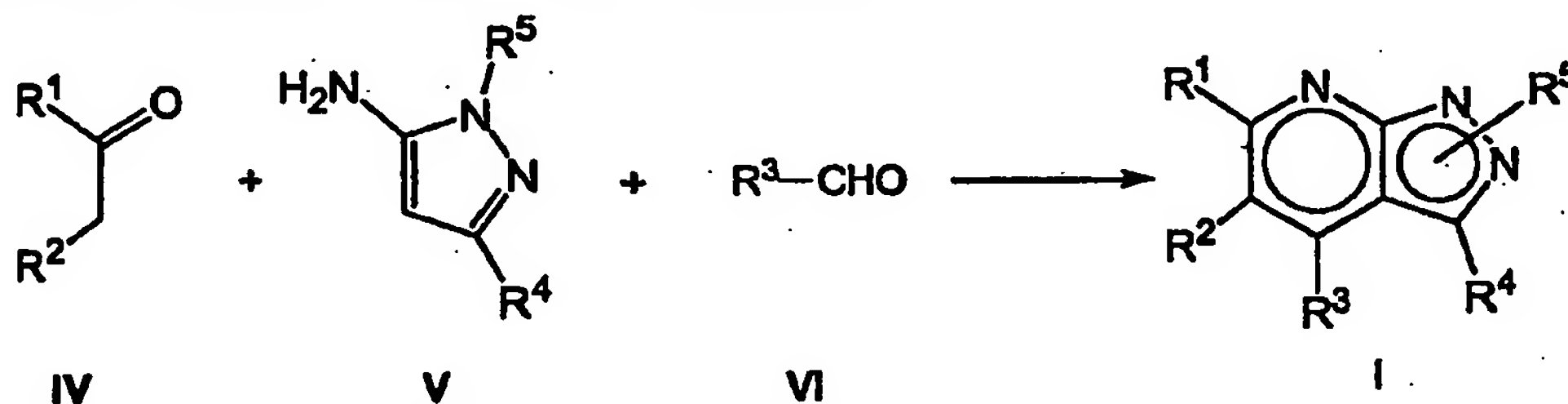
Some of the compounds of the present invention may exist as several diastereoisomers and/or several optical isomers. Diastereoisomers can be separated by conventional techniques such as chromatography or fractional
15 crystallization. Optical isomers can be resolved by conventional techniques of optical resolution to give optically pure isomers. This resolution can be carried out on any chiral synthetic intermediate or on products of general formula I. Optically pure isomers can also be individually obtained using enantiospecific synthesis. The present invention covers all isomers and mixtures thereof (for example
20 racemic mixtures) whether obtained by synthesis and also by physically mixing them.

Moreover, some compounds of the present invention may exhibit cis/trans isomers. The present invention includes each of the geometric isomers and its mixtures.

25 The compounds of formula I can be obtained by following the processes described below. As it is obvious to one skilled in the art the exact method used to prepare a given compound can vary depending on its chemical structure. Moreover, in some of the processes described below it may be necessary or advisable to protect the reactive or labile groups by conventional protective
30 groups. Both the nature of these protective groups and the procedures followed to introduce or remove these are well known and form part of the state of the art (see for example Greene T.W. and Wuts P.G.M, "Protective Groups in Organic Synthesis", John Wiley & Sons, 3rd edition, 1999). As an example, as protective

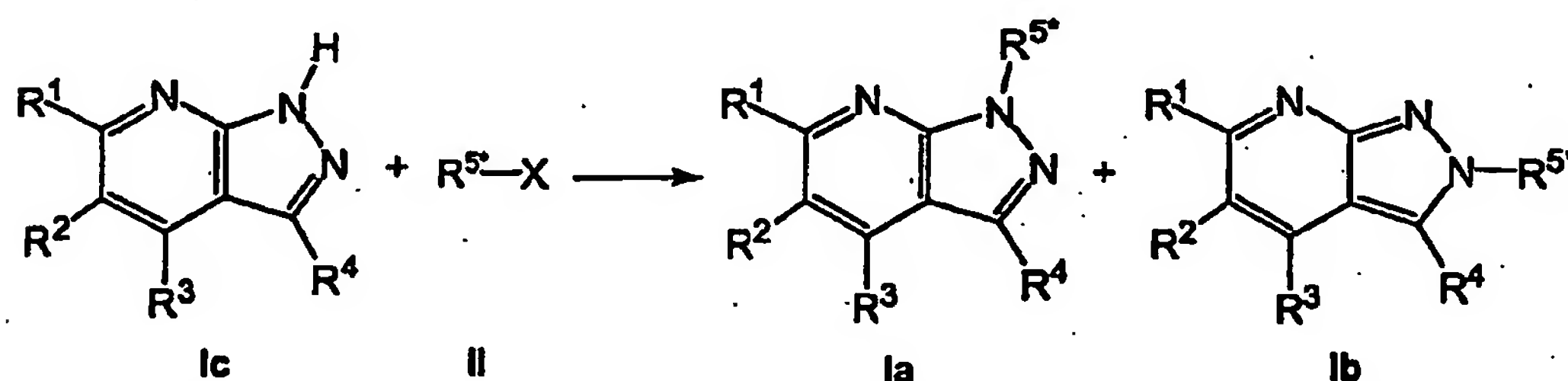
groups of an amino function *tert*-butoxycarbonyl (Boc) or benzyl (Bn) can be used. The carboxyl groups can be protected for example in the form of C₁₋₆alkyl esters or arylalkyl esters, such as benzyl, while the hydroxyl groups can be protected for example with tetrahydropyranyl (THP). Whenever a protective group is present a later deprotection step will be required, which can be performed under standard conditions in organic synthesis, such as those described in the above-mentioned reference.

The compounds of formula I can be obtained in general by reaction of a compound of formula IV with an aminopyrazole of formula V and an aldehyde of formula VI, as shown in the following scheme:



wherein R^1 , R^2 , R^3 , R^4 and R^5 have the same meaning as in general formula I. This reaction is carried out preferably in the presence of an inorganic acid such as hydrochloric acid, in a suitable polar solvent such as 2-methoxyethanol or ethanol, and heating, preferably to reflux.

The compounds of formula I wherein R^5 represents R^{5*} (i.e. compounds of formula Ia and Ib) are generally obtained by reaction of a compound of formula I wherein R^5 represents H (i.e. a compound of formula Ic) with an alkylating agent of formula $\text{R}^{5*}\text{-X}$ (II), as shown in the following scheme:



wherein R^{5*} represents R^a or R^f . R^1 , R^2 , R^3 , R^4 , R^a and R^f have the meaning described in general formula I and X represents a leaving group, for example an alkylsulfonate or an arylsulfonate such as mesylate or tosylate, or a halogen such as Cl, Br or I. This reaction is performed in the presence of a base such as KOH,

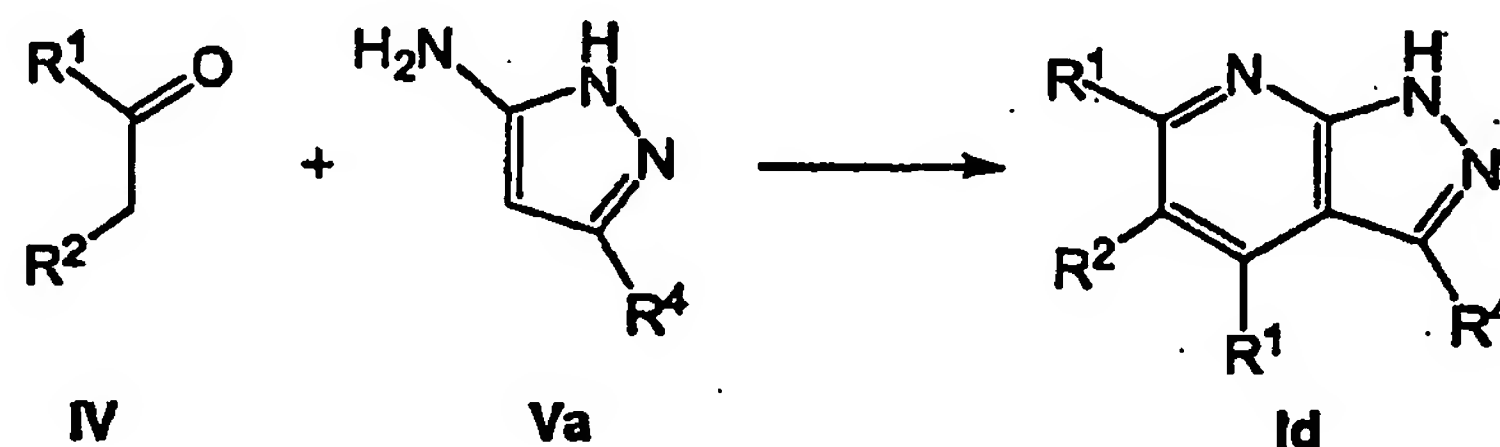
K₂CO₃ or NaH in a suitable solvent, such as for example acetone, toluene, dimethylformamide, 1,2-dimethoxyethane or diglyme, and heating, preferably to reflux. In the case of using apolar solvents such as toluene this reaction can be carried out in the presence of a cation sequestering agent such as crown ether 18-C-6 or a phase transfer agent such as a tetraalkylammonium salt.

Compounds of formula Ia and Ib wherein R^a represents -CH₂OH can be prepared by the reaction of a compound of formula Ic with formaldehyde in a suitable polar solvent, such as water, and heating, preferably to reflux.

Compounds of formula Ia and Ib wherein R^a represents optionally substituted phenyl or optionally substituted heteroaryl can be prepared by reaction of a compound of formula Ic with a boronic acid of formula R^a-B(OH)₂ (III) wherein R^a represents optionally substituted phenyl or heteroaryl in the presence of a base, such as pyridine and/or triethylamine, and in the presence of a catalyst, such as copper acetate (II), in a suitable solvent, such as an aprotic solvent such as dichloromethane.

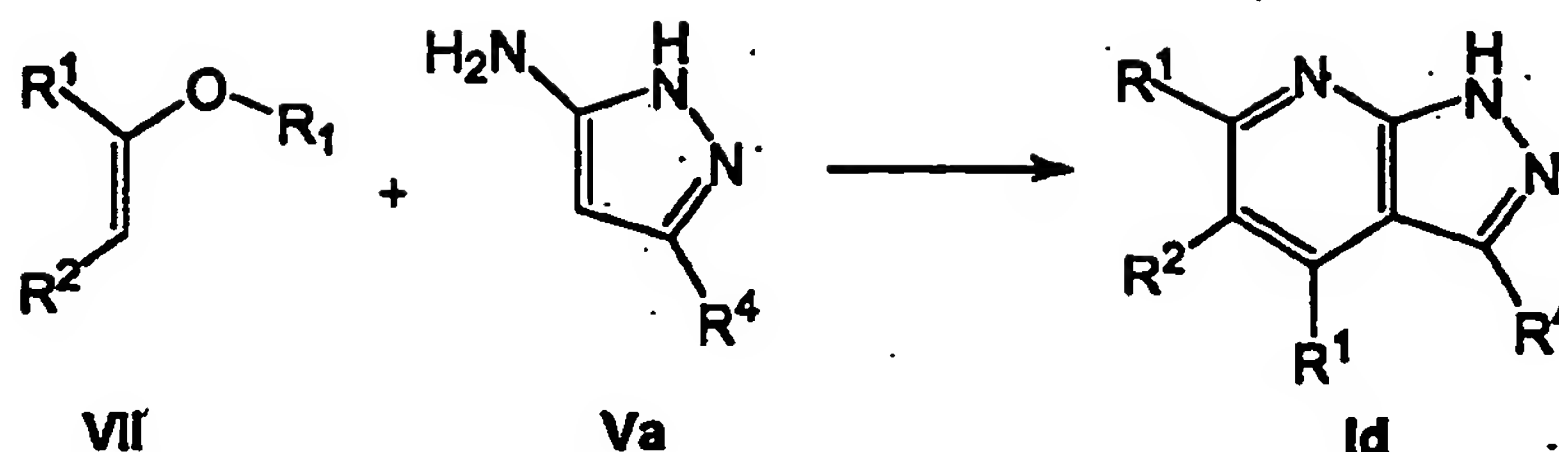
Compounds of formula II and III are commercially available, are widely described in the literature or can be prepared by methods similar to those described, and can be conveniently protected as required.

Compounds of formula I wherein R⁵ represents H and R³ has the same meaning as R¹ (i.e., a compound Id) can be also prepared by reaction of a compound of formula IV with an aminopyrazole of formula Va, as shown in the following scheme:



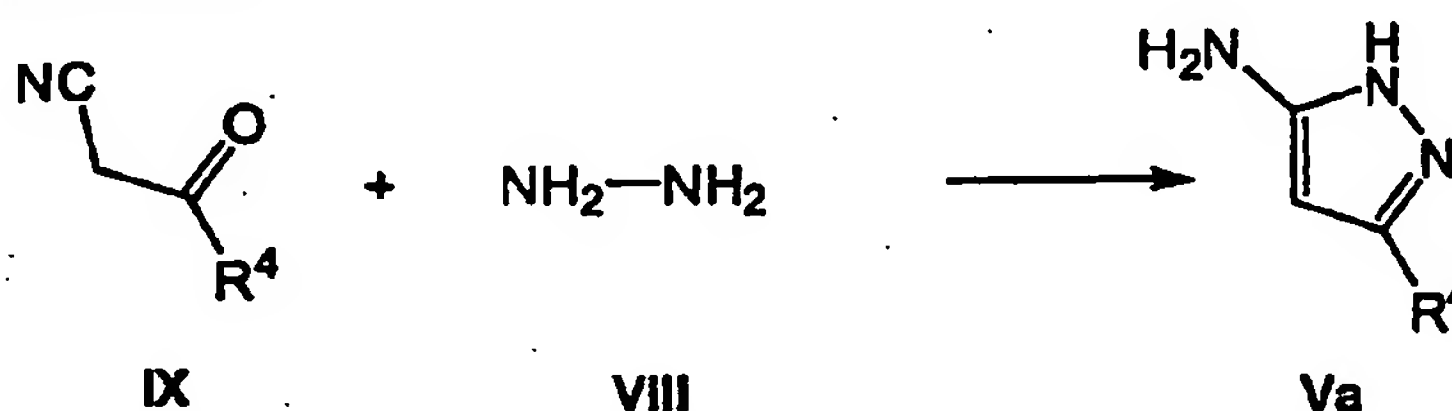
or alternatively by reaction of a compound of formula VII with a compound of formula Va

20



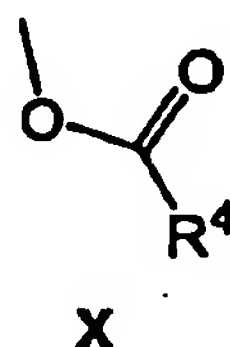
wherein R¹, R² and R⁴ have the meaning described in general formula I. This reaction is carried out preferably in the presence of an inorganic acid such as hydrochloric acid, in a suitable polar solvent such as 2-methoxyethanol or ethanol, and heating, preferably to reflux.

Aminopyrazoles of formula V and Va and aldehydes of formula VI, are commercially available and can be conveniently protected. Alternatively, compounds of formula Va can be conveniently prepared by reaction of a compound of formula VIII with a compound of formula IX, as shown in the following scheme:



wherein R⁴ has the meaning described in general formula I. This reaction is carried out in a suitable polar solvent such as ethanol and by heating, preferably to reflux. Compounds of formula V can also be obtained by this method, starting from a compound of formula IX and a compound of formula NH₂-NHR⁵ (VIIIa).

Compounds of formula IX are commercially available and can be conveniently protected or can be conveniently prepared by reaction of a compound of formula X



with acetonitrile, in the presence of a base such as butyl lithium, in a suitable solvent such as tetrahydrofuran and at a suitable temperature, preferably - 78°C.

Esters of formula X are commercially available or can be prepared by

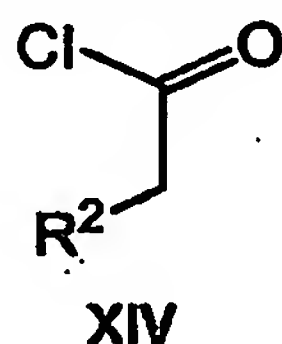
methods widely described in the literature and can be conveniently protected.

Enol ethers of formula VII can be conveniently prepared by reaction of a ketone of formula IV with a compound of formula $R^1\text{-COY}$ (XI) wherein Y represents a halogen, preferably Cl, in the presence of a base, such as NaH, in a suitable polar solvent such as dimethylformamide.

Compounds of formula XI are commercially available or can be prepared by conventional reactions from the corresponding carboxylic acids of formula $R^1\text{-CO}_2\text{H}$ (XII).

Compounds of formula XII are commercially available or can be prepared by methods widely described in the literature and can be conveniently protected.

Compounds of formula IV can be prepared by reaction of a compound of formula $R^1\text{-H}$ (XIII) with a compound of formula XIV



wherein R^1 and R^2 have the meaning described in general formula I, in the presence of a Lewis acid, such as AlCl_3 , in a suitable halogenated solvent such as dichloromethane.

Compounds of formula XIV are commercially available or can be prepared readily from the corresponding carboxylic acid following conventional procedures.

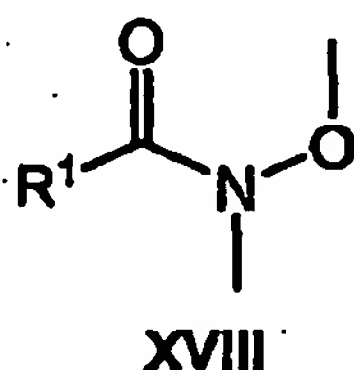
Alternatively, compounds of formula IV can be conveniently prepared by reaction of a compound of formula $R^2\text{-CH}_3$ (XV) with a compound of formula $R^1\text{-CN}$ (XVI), wherein R^1 and R^2 have the meaning described in general formula I, in the presence of a base such as lithium diisopropylamide, generated from butyl lithium and *N,N'*-diisopropylamine, in an aprotic polar solvent, such as tetrahydrofuran and cooling, preferably to -78°C .

Alternatively, compounds of formula IV can be conveniently prepared by reaction of a compound of formula $R^2\text{-CH}_3$ (XV) with a compound of formula $R^1\text{-CO}_2\text{R}^6$ (XVII), wherein R^1 and R^2 have the meaning described in general formula I and R^6 represents C_{1-6} alkyl, in the presence of a base such as sodium bis(trimethylsilyl)amide or sodium hydride, in an aprotic polar solvent, such as tetrahydrofuran or dimethylformamide and at a suitable temperature, preferably

room temperature.

The compounds XIII, XV, XVI and XVII are commercially available or can be prepared by methods widely described in the literature.

Alternatively, compounds of formula IV can be conveniently prepared by
5 reaction of a compound of formula R^2-CH_3 (XV) with a compound of formula XVIII in the same conditions as those described previously for reaction of the compound of formula XV with a compound of formula XVI.

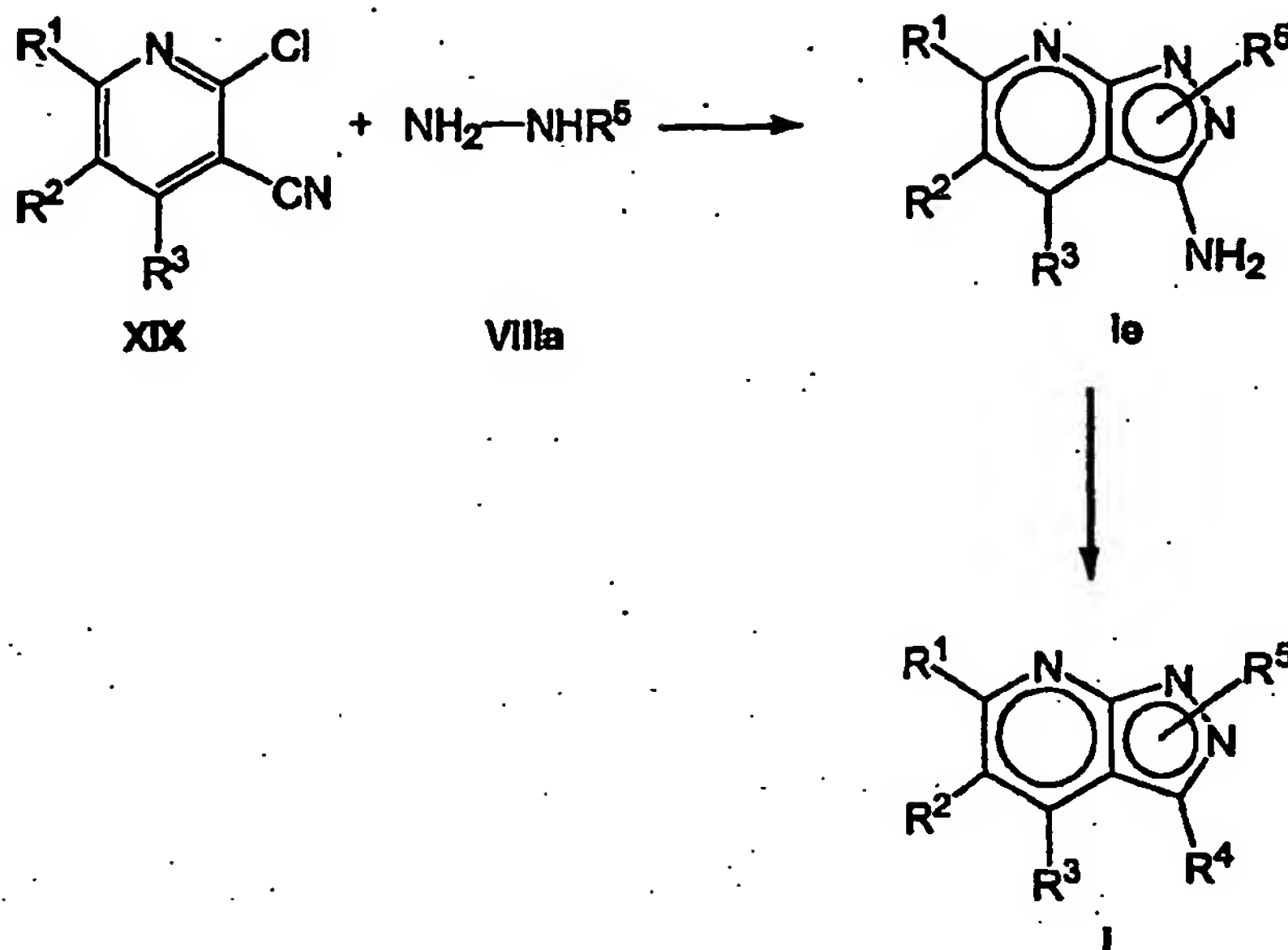


Compounds of formula XVIII can be conveniently prepared by reaction of a
10 compound of formula XI with *N,O*-dimethylhydroxylamine hydrochloride in the presence of a base, such as triethylamine, in a suitable halogenated solvent such as dichloromethane and cooling, preferably to 0 °C.

Alternatively, the compounds of formula XVIII can be prepared by reaction
of a compound of formula XII with *N,O*-dimethylhydroxylamine hydrochloride in the
15 presence of a suitable condensing agent such as for example *N*-(3-dimethylaminopropyl)-*N'*-ethylcarbodiimide or dicyclohexylcarbodiimide optionally in the presence of 1-hydroxybenzotriazole, or in the presence of a suitable base, such as pyridine, in a suitable solvent such as dimethylformamide.

The compounds of formula I can also be obtained from a compound of
20 formula XIX, as shown in the scheme below.

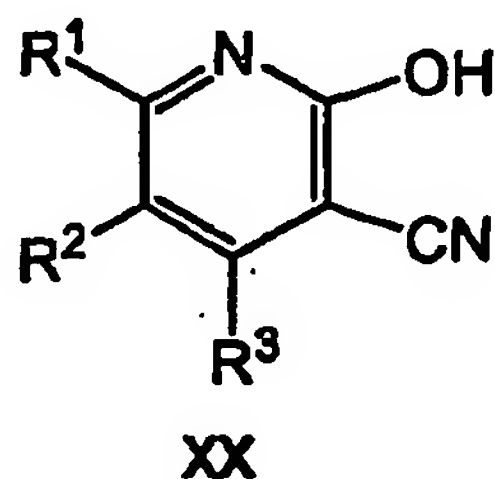
23



wherein R¹, R², R³, R⁴ and R⁵ have the meaning described in general formula I. The reaction of XIX with a compound of formula VIIIa gives rise to a compound of formula I wherein R⁴ = NH₂ (Ie). This reaction is carried out in a suitable solvent such as ethanol and heating, preferably to reflux. Starting from these compounds of formula I wherein R⁴ = NH₂, compounds of formula I wherein R⁴ is different from NH₂ can be generated by interconversion reactions, explained in more detail further on. This method is useful to prepare compounds of formula I wherein R³ represents H, optionally substituted C₁₋₆alkyl or Cy different from R¹.

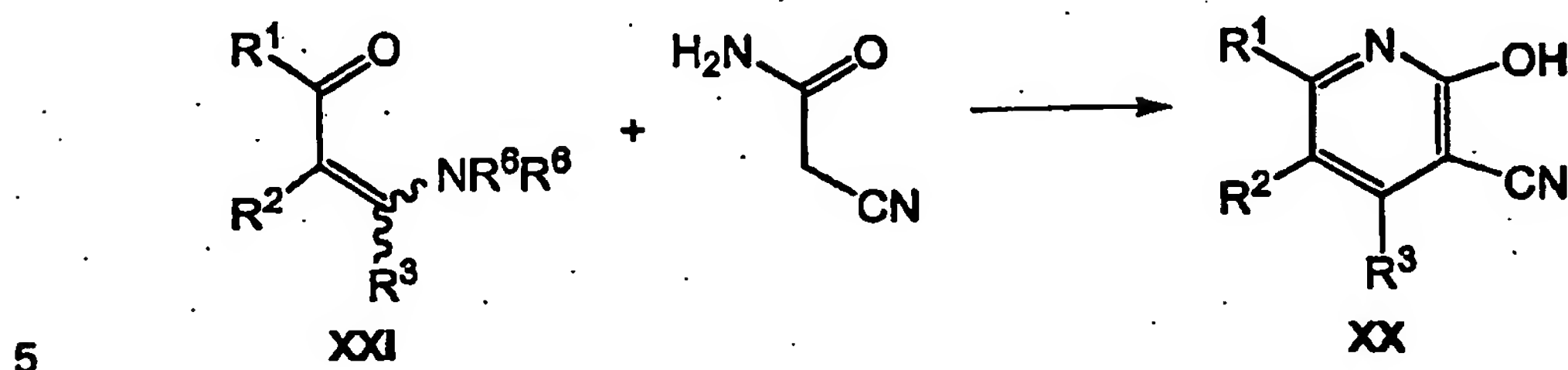
Compounds of formula VIIIa are commercially available and can be conveniently protected.

Compounds of formula XIX can be conveniently prepared by reaction of a compound of formula XX



wherein R¹, R² and R³ have the meaning described in general formula I, with a chlorinating agent such as POCl₃ or PCl₃ without solvent or in a suitable solvent such as dimethylformamide and heating, preferably to reflux.

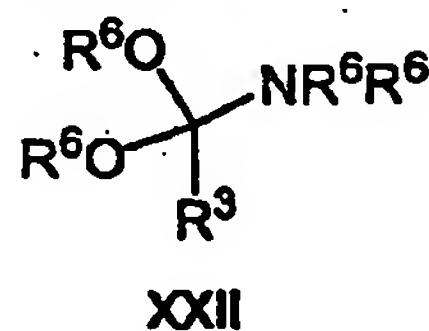
Compounds of formula **XX** are generally obtained by reaction of a compound of formula **XXI** with 2-cyanoacetamide, as shown in the following scheme:



wherein R^1 , R^2 and R^3 have the meaning described in general formula I and each R^6 independently represents C_{1-6} alkyl. This reaction is carried out in the presence of a base such as sodium methoxide, in a suitable solvent such as dimethylformamide and heating, preferably to reflux.

10

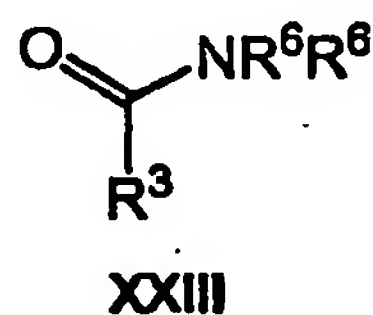
Compounds of formula **XXI** can be conveniently prepared by reaction of a compound of formula **IV** with a compound of formula **XXII**



wherein R^3 and R^6 have the meaning previously described, in a suitable solvent such as tetrahydrofuran.

15

Compounds of formula **XXII** are commercially available or can be prepared by methods described in the literature, for example by reacting an amide of formula **XXIII**



20 with triethyloxonium tetrafluoroborate in the presence of a base such as sodium ethoxide in a suitable solvent such as ethanol or a mixture of ethanol-diethyl ether.

Alternatively, compounds of the present invention can also be obtained by interconversion of another compound of formula I, in one or several steps, by using well-known reactions in organic chemistry under the reported standard experimental conditions.

25

Thus, R^4 can be transformed into another R^4 , generating new compounds of formula I. For example, $R^4 = H$ can be transformed into $R^4 = Br$ by reaction with an appropriate brominating agent, such as Br_2 , in a suitable solvent such as chloroform, and at an appropriate temperature comprised between room
5 temperature and the boiling point of the solvent;

or $R^4 = H$ can be converted into $R^4 = Cl$ by reaction with an appropriate chlorinating agent such as *N*-chlorosuccinimide, in a suitable solvent such as dimethylformamide at an appropriate temperature comprised between room temperature and the boiling point of the solvent;

10 or $R^4 = NH_2$ can be converted into $R^4 = \text{halogen}$ by formation of a diazonium salt with $NaNO_2$ followed by reaction with a copper halide such as $CuBr$ or $CuCl$, in the presence of an acid such as for example HBr or HCl ;

or $R^4 = NH_2$ can be converted into $R^4 = H$ by forming a diazonium salt with $NaNO_2$ followed by reaction with H_3PO_2 , in a suitable solvent such as water;

15 or $R^4 = \text{halogen}$ can be converted into $R^4 = CN$ by reaction with a cyanide salt such as $CuCN$, in a suitable solvent such as *N*-methylpyrrolidone and heating, preferably to reflux.

Other conversions of R^4 , which can also be applied to R^5 and also to the substituents of the R^1 , R^2 and R^3 groups to generate other compounds of formula I
20 include, for example:

conversion of CN into $CONH_2$ by hydrolysis with a base such as KOH in a suitable solvent such as *tert*-butanol and heating, preferably to reflux;

conversion of CN into CH_2NH_2 by reaction with a reducing agent, such as $LiAlH_4$, in a suitable solvent such as diethyl ether;

25 conversion of a carboxylic acid group into an ester or an amide by reaction with an alcohol or an amine, respectively, in the presence of an activating agent such as *N,N'*-dicyclohexylcarbodiimide and 4-dimethylaminopyridine and in a suitable solvent such as diethyl ether;

conversion of an ester group into a carboxylic group by hydrolysis in the
30 presence of a base, such as KOH , in a suitable solvent such as ethanol;

conversion of OH , SH or NH_2 into OR , SR and NHR or NRR , respectively, by reaction of an alkylating agent $R-X$ wherein R represents R^a , R^c , R^e or R^f and X represents halogen, preferably chloro or bromo, in the presence of a base such as

triethylamine, sodium hydroxide, sodium carbonate, potassium carbonate or sodium hydride in a suitable solvent such as dichloromethane, chloroform, dimethylformamide, ethanol or butanol, at a temperature comprised between room temperature and the temperature of the boiling point of the solvent;

- 5 alternatively, NHR can be converted into NCH_3R , wherein R represents R^a , R^c , R^e or R^f , by reaction with formaldehyde in acid medium, such as formic acid and heating, preferably to reflux;

- conversion of an amine into an amide by reaction of a carboxylic acid in the presence of an appropriate condensing agent such as *N*-(3-dimethylaminopropyl)-*N'*-ethylcarbodiimide or dicyclohexylcarbodiimide optionally in the presence of 1-
10 hydroxybenzotriazole, or in the presence of an appropriate base such as pyridine, in a suitable solvent, such as dimethylformamide; or alternatively, an amine can be converted into an amide by reaction with an acyl chloride in pyridine or in the presence of a base such as triethylamine in a suitable solvent such as
15 dichloromethane, and cooling, preferably to 0 °C;

- conversion of an amine into a urea or a carbamate by a two step sequence that involves reacting the amine with an activating agent such as triphosgene, in the presence of a base such as diisopropylethylamine, triethylamine or *N*-methylmorpholine, in a suitable solvent such as acetonitrile or a halogenated
20 hydrocarbon such as chloroform or dichloromethane, and then reacting the resulting compound with the second amine in the case of a urea or with an alcohol in the case of a carbamate, in a suitable solvent, such as the solvent used in the first step; or alternatively an amine can be converted into a urea or carbamate by reaction with an isocyanate or a chloroformate, respectively, in a suitable solvent,
25 such as dimethylformamide, at a suitable temperature, preferably room temperature;

- conversion of an amine into a sulfonamide by reaction with a sulfonyl halide, such as sulfonyl chloride, optionally in the presence of a base such as 4-dimethylaminopyridine, in a suitable solvent such as dioxane, chloroform,
30 dichloromethane or pyridine;

conversion of a hydroxyl group into an ester group by reaction with a carboxylic acid in the standard conditions mentioned previously;

conversion of a sulfanyl group into a sulfinyl or sulfonyl group by reaction

with 1 or 2 equivalents, respectively, of an appropriate oxidizing agent such as *m*-chloroperbenzoic acid in a suitable solvent such as for example dichloromethane;

conversion of a primary or secondary hydroxyl group into a leaving group, for example an alkylsulfonate or arylsulfonate such as mesylate or tosylate or a
5 halogen such as Cl, Br or I, by reaction with a sulfonyl halide such as methanesulfonyl chloride, in the presence of a base, such as pyridine or triethylamine, in a suitable solvent such as dichloromethane or chloroform, or with a halogenating agent, such as SOCl₂, in a suitable solvent such as tetrahydrofuran;

10 the substitution of said leaving group by reaction with an alcohol, amine or thiol, optionally in the presence of a base, such as K₂CO₃, and in a suitable solvent such as dimethylformamide, 1,2-dimethoxyethane or acetonitrile;

the elimination of a leaving group bonded to an alkyl group to give an alkenyl group by reaction with a base, such as KOH, in a suitable solvent, such as
15 toluene and heating, preferably to reflux;

conversion of a primary amide into a secondary amide by reaction with an alkylating agent in the presence of a strong base, such as sodium hydride, in a suitable solvent and at a temperature comprised between room temperature and the boiling point of the solvent;

20 conversion of CHO into an amine by reaction with an amine in the presence of a reducing agent such as sodium triacetoxyborohydride, in a suitable solvent such as for example 1,2-dichloroethane;

conversion of an acetal group into an aldehyde by reaction in acid medium, for example in HCl, at an appropriate temperature, preferably to reflux;

25 conversion of an ester group into an alcohol group by reaction with a reducing agent, such as LiAlH₄, in a suitable solvent, such as tetrahydrofuran;

conversion of a sulfonyl group bonded to an aromatic ring into an amino derivative by reaction with the corresponding amine, in a suitable solvent such as tetrahydrofuran or using the amine as a solvent, heating, preferably to a
30 temperature comprised between room temperature and 100 °C and preferably carrying out the reaction in a closed vessel under atmospheric pressure;

conversion of a sulfonyl group bonded to an aromatic ring by displacement with an alkoxide to give the corresponding alkoxy derivative, in a suitable solvent

such as tetrahydrofuran or using the corresponding alcohol as a solvent, heating, preferably to a temperature comprised between room temperature and 100 °C and preferably carrying out the reaction in a closed vessel under atmospheric pressure;

5 conversion of a vinylic or aromatic halogen into NHR, wherein R represents R^a, R^c, R^e or R^f, by reacting with an amine of formula H₂NR and preferably heating;

alternatively, a vinylic or aromatic halogen can be converted into NHR by reaction with an amine of formula H₂NR in the presence of a base, such as
10 Cs₂CO₃ or sodium *tert*-butoxide, in the presence of a palladium catalyst, such as palladium acetate (II), and a phosphine such as 2,2'-bis(diphenylphosphino)-1,1'-binaphthyl, in a solvent, such as toluene, and preferably heating;

conversion of a vinylic or aromatic halogen into a phenyl or heteroaryl group by treatment with a phenyl- or a heteroarylboronic acid in the presence of a
15 catalyst, such as a palladium catalyst, such as palladium acetate (II) or tetraakis(triphenylphosphine) palladium (0) and of a base such as Na₂CO₃, K₂CO₃ or CsF, in a suitable polar solvent, such as 1,2-dimethoxyethane or toluene-water mixtures, and preferably heating;

conversion of an aromatic halogen into H by halogenolysis, with a reducing
20 agent such as Zn, in a suitable solvent, such as acetic acid and heating, preferably to reflux; and

oxidation of the N at position 7 of the pyrazolo[3,4-*b*]pyridine ring to give the corresponding N-oxide by reaction with an oxidizing agent, such as *m*-chloroperbenzoic acid, in a suitable solvent, such as for example dichloromethane.

25 Likewise, any of the aromatic rings of the compounds of the present invention can undergo electrophilic aromatic substitution reactions, widely described in the literature.

Many of these interconversion reactions are explained in greater detail in the examples.

30 As it will be obvious to those skilled in the art, these interconversion reactions can also be carried out on synthesis intermediates of compounds of formula I.

The present invention also includes the salts of the compounds of formula I.

These salts can be prepared by conventional methods by treating a compound of formula I with an appropriate amount of an acid such as hydrochloric acid, sulfuric acid, nitric acid, oxalic acid or methanesulfonic acid. In the case of compounds of formula I that contain an acidic proton, salts can also be obtained by treatment
5 with a base such as sodium hydroxide, potassium hydroxide, calcium hydroxide or calcium carbonate. Salts of the compounds of formula I can be converted in turn into other salts of compounds of formula I by ion exchange using a ionic exchange resin.

As mentioned previously, the compounds of the present invention act as
10 p38 kinase inhibitors, inducing reduction of proinflammatory cytokines. Therefore, these compounds are expected to be useful to treat or prevent diseases in which kinase p38 plays a role. This includes diseases caused by overproduction of cytokines such as TNF- α , IL-1, IL-6 or IL-8. These diseases include, but are not limited to, immune, autoimmune and inflammatory diseases, cardiovascular
15 diseases, infectious diseases, bone resorption disorders, neurodegenerative diseases, proliferative diseases and processes associated with cyclooxygenase-2 induction.

As an example, immune, autoimmune and inflammatory diseases that can be treated or prevented with the compounds of the present invention include
20 rheumatic diseases (e.g. rheumatoid arthritis, psoriatic arthritis, infectious arthritis, progressive chronic arthritis, deforming arthritis, osteoarthritis, traumatic arthritis, gouty arthritis, Reiter's syndrome, polychondritis, acute synovitis and spondylitis), glomerulonephritis (with or without nephrotic syndrome), autoimmune hematologic disorders (e.g. hemolytic anemia, aplasic anemia, idiopathic thrombocytopenia
25 and neutropenia), autoimmune gastritis and autoimmune inflammatory bowel diseases (e.g. ulcerative colitis and Crohn's disease), host versus graft disease, allograft rejection, chronic thyroiditis, Graves' disease, scleroderma, diabetes (type I and type II), active hepatitis (acute and chronic), primary biliary cirrhosis, myasthenia gravis, multiple sclerosis, systemic lupus erythematosus, psoriasis,
30 atopic dermatitis, contact dermatitis, eczema, skin sunburns, chronic renal insufficiency, Stevens-Johnson syndrome, idiopathic sprue, sarcoidosis, Guillain-Barré syndrome, uveitis, conjunctivitis, keratoconjunctivitis, otitis media, periodontal disease, pulmonary interstitial fibrosis, asthma, bronchitis, rhinitis,

sinusitis, pneumoconiosis, pulmonary insufficiency syndrome, pulmonary emphysema, pulmonary fibrosis, silicosis, chronic inflammatory pulmonary disease (e.g. chronic obstructive pulmonary disease) and other inflammatory or obstructive diseases of the airways.

5 Cardiovascular diseases that can be treated or prevented include, among others, myocardial infarction, cardiac hypertrophy, cardiac insufficiency, ischaemia-reperfusion disorders, thrombosis, thrombin-induced platelet aggregation, acute coronary syndromes, atherosclerosis and cerebrovascular accidents.

10 Infectious diseases that can be treated or prevented include, among others, sepsis, septic shock, endotoxic shock, sepsis by Gram-negative bacteria, shigellosis, meningitis, cerebral malaria, pneumonia, tuberculosis, viral myocarditis, viral hepatitis (hepatitis A, hepatitis B and hepatitis C), HIV infection, retinitis caused by cytomegalovirus, influenza, herpes, treatment of infections
15 associated with severe burns, myalgias caused by infections, cachexia secondary to infections, and veterinary viral infections such as lentivirus, caprine arthritis virus, visna-maedi virus, feline immunodeficiency virus, bovine immunodeficiency virus or canine immunodeficiency virus.

Bone resorption disorders that can be treated or prevented include
20 osteoporosis, osteoarthritis, traumatic arthritis, gouty arthritis and bone disorders related with multiple myeloma, among others.

Neurodegenerative diseases that can be treated or prevented include Alzheimer's disease, Parkinson's disease, cerebral ischaemia and traumatic neurodegenerative disease, among others.

25 Proliferative diseases that can be treated or prevented include endometriosis, solid tumors, acute and chronic myeloid leukemia, Kaposi sarcoma, multiple myeloma, metastatic melanoma and angioendothelial disorders such as ocular neovascularisation and infantile haemangioma.

p38 kinase inhibitors also inhibit the expression of proinflammatory proteins
30 such as cyclooxygenase-2 (COX-2), the enzyme responsible for prostaglandin production. Therefore, the compounds of the present invention can also be used to treat or prevent diseases mediated by COX-2 and especially to treat processes

with inflammation, fever and neuromuscular pain such as cephalaea, pain caused by cancer, tooth pain and arthritic pain.

According to the activity of the products herein described, the present invention also relates to compositions which contain a compound of the present invention, together with an excipient or other auxiliary agents if necessary. The compounds of the present invention can be administered in the form of any pharmaceutical formulation, the nature of which, as it is well known, will depend upon the nature of the active compound and its route of administration. Any route of administration may be used, for example oral, parenteral, nasal, ocular, rectal and topical administration.

According to the present invention, solid compositions for oral administration include tablets, granulates and capsules. In any case the manufacturing method is based on a simple mixture, dry granulation or wet granulation of the active compound with excipients. These excipients can be, for example, diluents such as lactose, microcrystalline cellulose, mannitol or calcium hydrogenphosphate; binding agents such as for example starch, gelatin or povidone; disintegrants such as sodium carboxymethyl starch or sodium croscarmellose; and lubricating agents such as for example magnesium stearate, stearic acid or talc. Tablets can be additionally coated with suitable excipients by using known techniques with the purpose of delaying their disintegration and absorption in the gastrointestinal tract and thereby provide a sustained action over a longer period, or simply to improve their organoleptic properties or their stability. The active compound can also be incorporated by coating onto inert *pellets* using natural or synthetic film-coating agents. Soft gelatin capsules are also possible, in which the active compound is mixed with water or an oily medium, for example coconut oil, mineral oil or olive oil.

Powders and granulates for the preparation of oral suspensions by the addition of water can be obtained by mixing the active compound with dispersing or wetting agents; suspending agents and preservatives. Other excipients can also be added, for example sweetening, flavouring and colouring agents.

Liquid forms for oral administration include emulsions, solutions, suspensions, syrups and elixirs containing commonly-used inert diluents, such as purified water, ethanol, sorbitol, glycerol, polyethylene glycols (macrogols) and

propylene glycol. Said compositions can also contain coadjuvants such as wetting, suspending, sweetening, flavouring agents, preservatives and buffers.

Injectable preparations, according to the present invention, for parenteral administration, comprise sterile solutions, suspensions or emulsions, in an aqueous or non-aqueous solvent such as propylene glycol, polyethylene glycol or vegetable oils. These compositions can also contain coadjuvants, such as wetting, emulsifying, dispersing agents and preservatives. They may be sterilized by any known method or prepared as sterile solid compositions which will be dissolved in water or any other sterile injectable medium immediately before use. It is also possible to start from sterile materials and keep them under these conditions throughout all the manufacturing process.

For the rectal administration, the active compound can be preferably formulated as a suppository on an oily base, such as for example vegetable oils or solid semisynthetic glycerides, or on a hydrophilic base such as polyethylene glycols (macrogol).

The compound can also be formulated for its topical application for the treatment of pathologies occurring in zones or organs accessible through this route, such as eyes, skin and the intestinal tract. Formulations include creams, lotions, gels, powders, solutions and patches wherein the compound is dispersed or dissolved in suitable excipients.

The compounds of the present invention can also be formulated as a solid form, dissolved or dispersed in a suitable vehicle, for inhalation in single or multidose container. Preparations to be administered as an aerosol (dispersion of solid or liquid particles in a gas) use suitable devices such as nebulisers, pressured metered-dose inhalers or dry-powder inhalers. Depending on this, the compound will be formulated with excipients such as propellants responsible for developing the proper pressure within the container to force the content out through the opening of the valve, solvents, emulsifying agents, viscosity-increasing agents, preservatives, stabilizing agents and lubricants to avoid the blockade of the valve.

The dosage and frequency of doses will depend upon the nature and severity of the disease to be treated, the age, the general condition and body weight of the patient, as well as the particular compound

administered and the route of administration, among other factors. A representative example of a suitable dosage range is from about 0.01 mg/Kg to about 100 mg/Kg per day, which can be administered as a single or divided doses. However, the dosage administered is generally left to the discretion of the physician.

The activity of the compounds of this invention can be assessed using the following tests:

Test 1: Inhibition of TNF- α release induced by LPS in human histiocytic lymphoma cells, U-937

Maintenance and differentiation of U-937 cells: U-937 cells (ATCC N° CRL-159.2) are cultivated in RPMI 1640 medium supplemented with 10% inactivated fetal bovine serum (Gibco). A total of 0.5×10^6 cells are incubated in the presence of 20 ng/mL of PMA (*phorbol 12-myristate 13-acetate*) for 24 h to achieve complete monocytic differentiation. All the incubations are carried out at 37°C in an atmosphere with 5% CO₂. The cells are centrifuged (200 x g for 5 min) and resuspended in RPMI 1640 medium supplemented with 2% inactivated fetal bovine serum at a density of 2×10^6 cells/mL.

Inhibition of TNF- α release: 100 μ L of cells U-937 (2×10^6 cells/mL) are incubated with 100 μ L of the test product (final concentration, 0.001-10 μ M) for 30 min in 96- well plates. The mother solutions of the products (10 mM in DMSO) are diluted in culture medium to reach a final DMSO concentration equal to or less than 0.1 %. A total of 20 μ L of LPS (*E. coli* 055B5, Sigma) are added to a final concentration of 100 ng/mL and after incubation for 4 hours the amount of TNF- α released in the supernatant is quantified using a commercial ELISA kit (Biosource International).

Test 2: Inhibition of TNF- α release induced by LPS in human peripheral blood mononuclear cells

To obtain the mononuclear cells: heparinized venous blood, obtained from healthy volunteers, is diluted with an equal volume of saline phosphate buffer without calcium or magnesium. Aliquots of 30 mL of the mixture are transferred to 50 mL centrifuge tubes containing 15 mL of Ficoll-Hypaque (1.077 g/mL). The tubes are

centrifuged at 1200 x g for 20 min at room temperature without braking. Approximately two-thirds of the band of platelets lying above the mononuclear cells is removed with a pipette. The mononuclear cells are carefully transferred to a 50 mL tube, washed twice with saline phosphate buffer, centrifuged at 300 x g
5 for 10 min at room temperature and resuspended in RPMI supplemented with 1% inactivated fetal bovine serum at a cell density of 2×10^6 cells/mL.

Inhibition of TNF- α release: 100 μ L of mononuclear cells (2×10^6 cells/mL) are incubated on 96-well plates with 50 μ L of the test product (final concentration, 0.001-10 μ M) and 50 μ L LPS (*E. coli* 055B5, Sigma) at a final concentration of 400
10 ng/mL for 19 h at 37 °C in an atmosphere with CO₂ at 5%. The amount of TNF- α released in the supernatant is quantified using a commercial ELISA kit (Biosource International).

Test 3: Inhibition of p38- α kinase:

15 In a final volume of 25 μ L, a total of 5 μ L of the test product (final concentration, 0.001-10 μ M), 5-10 mU of p38- α with 0.33 mg/mL of myelin basic protein, Mg²⁺ acetate (10 mM) and [γ -³³P-ATP] (100 μ M, specific activity 500 cpm/pmol) in buffer Tris 25 mM pH7.5, EGTA 0.02 mM is incubated. The reaction is started by adding Mg²⁺[γ -³³P-ATP]. After incubation for 40 min at room temperature, the reaction is
20 quenched by adding 5 μ L of 3% phosphoric acid solution. The reaction mixture (10 μ L) is passed through a filter (P30) and washed three times for 5 min with a 75 mM phosphoric acid solution and once with methanol before drying it and counting it, by liquid scintillation.

25 The following table shows the results obtained with representative compounds of the invention in test 2:

Example	% Inhibition at 0.1 μ M
1	57.5
4	68.4
6	81.4

8	68.3
18	82.6
22	56.5
30	83.6
36	92.0
39	51.1
41	90.6
43	61.2
56	58.7
59	60.0
63	53.2
68	50.0
72	52.7
78	73.3
80	69.3
82	59.9
90	86.3
102	67.9
106	69.3
121	52.1
128	82.0
136	67.0
137	61.4
183	68.7
184	71.2
188	73.2
196	70.2
208	67.7
209	84.2
210	57.3
211	70.6

212	67.5
213	68.1
214	71.1
217	53.8
232	67.9
237	53.4
240	58.4
248	53.2
250	67.0
268	69.1
272	65.6
279	100.0
282	65.2
283	52.4
287	71.2
289	51.3
290	65.8

The following examples illustrate, but do not limit, the scope of the invention.

The following abbreviations have been used in the examples:

- 5 AcOH: acetic acid
- EtOAc: ethyl acetate
- NH₄OAc: ammonium acetate
- BuLi: butyl lithium
- t*BuOH: *tert*-butanol
- 10 conc.: concentrated
- DMAP: 4-(*N,N*-dimethylamino)pyridine
- DMF: dimethylformamide
- EtOH: ethanol
- MeOH: methanol
- 15 THF: tetrahydrofuran

t_R : retention time

LC-MS: liquid chromatography-mass spectrometry

LC-MS spectra have been performed using the following chromatographic
5 methods:

Method 1: Column Tracer Excel 120, ODSB 5 μm (10 mm x 0.21 mm),
temperature: 30 °C, flow: 0.35 mL/min, eluent: A= acetonitrile, B = 0.1 % HCOOH,
gradient: 0 min 10% A - 10 min 90% A.

Method 2: Column X-Terra MS C18 5 μm (150 mm x 2.1 mm), temperature: 30 °C,
10 flow: 0.40 mL/min, eluent: A= acetonitrile, B = 10 mM NH_4OAc (pH = 6.80),
gradient: 0 min 25% A - 6 min 80% A - 7.5 min 25% A.

REFERENCE EXAMPLE 1

1-(4-Fluorophenyl)-2-(4-pyridyl)ethanone

15 a) 4-Fluoro-*N*-methoxy-*N*-methylbenzamide

In a volumetric flask *N,O*-dimethylhydroxylamine hydrochloride (25.54 g, 261.8
mmol) and CH_2Cl_2 (443 mL) were introduced under argon atmosphere at 0°C. 4-
Fluorobenzoyl chloride (34.59 g, 218.2 mmol) was added followed by the slow
addition of triethylamine (48.13 g, 475.6 mmol). The reaction was stirred for 30
20 min at 5 °C and allowed to reach room temperature. It was washed with 5%
aqueous citric acid (180 mL) and with 5% aqueous NaHCO_3 (180 mL). The
aqueous phase was extracted with CH_2Cl_2 . The organic phase was dried over
 Na_2SO_4 and concentrated to dryness, to afford 20.23 g of the desired compound
(yield: 88%).

25 b) Title compound

To a solution of diisopropylamine (23.4 mL, 165.7 mmol) in THF (250 mL), cooled
to -78 °C, BuLi (103.5 mL of a 1.6 M solution in hexane, 165.7 mmol) was added
dropwise under argon atmosphere. After 5 minutes, a solution of 4-methylpyridine
(10.28 g, 110.4 mmol) in THF (85 mL) was added over 20 min. The mixture was
30 stirred at 0°C for 15 min and a solution of 4-fluoro-*N*-methoxy-*N*-methylbenzamide
(obtained in section a) in THF (85 mL) was added over 30 min period. The
reaction was allowed to reach room temperature. Water (100 mL) and EtOAc (100
mL) were added and the mixture was stirred for 30 min. The organic phase was

separated, dried over Na_2SO_4 and concentrated to dryness, to afford 24.32 g of the desired compound (yield: 100%).

^1H NMR (300 MHz, CDCl_3) δ (TMS): 4.29 (s, 2 H), 7.14 – 7.23 (complex signal, 4 H), 8.05 (m, 2 H), 8.59 (dd, $J_o = 1.6$ Hz, $J_m = 4.4$ Hz, 2 H).

5

REFERENCE EXAMPLE 2

2-(4-Pyridyl)-1-[3-(trifluoromethyl)phenyl]ethanone

a) *N*-Methoxy-*N*-methyl-3-(trifluoromethyl)benzamide

Following a similar procedure to that described in reference example 1 section a, but using 3-(trifluoromethyl)benzoyl chloride instead of 4-fluorobenzoyl chloride, the desired product was obtained (yield: 86%).

10

b) Title compound

Following a similar procedure to that described in reference example 1 section b, but using *N*-methoxy-*N*-methyl-3-(trifluoromethyl)benzamide (obtained in section a of this example) instead of 4-fluoro-*N*-methoxy-*N*-methylbenzamide, the title compound was obtained (yield: 22%).

15

^1H NMR (300 MHz, CDCl_3) δ (TMS): 4.31 (s, 2 H), 7.20 (d, $J = 5.8$ Hz, 2 H), 7.63 (t, $J = 7.8$ Hz, 1 H), 7.84 (d, $J = 7.8$ Hz, 1 H), 8.16 (d, $J = 7.9$ Hz, 1 H), 8.24 (s, 1 H), 8.56 (d, $J = 5.8$ Hz, 2 H).

REFERENCE EXAMPLE 3

20

1-Phenyl-2-(4-pyridyl)ethanone

A solution of diisopropylamine (22 mL, 15.03 mmol) in THF (200 mL) under argon atmosphere was cooled to -78°C . BuLi (96 mL of a 1.6 M solution in hexane, 153.0 mmol) was added dropwise. After 1 h, a solution of 4-methylpyridine (15.00 g, 161.1 mmol) in THF (75 mL) was added and allowed to warm to 0°C . At this temperature it was stirred for 30 minutes. It was cooled to -78°C and benzonitrile (18.27 g, 177.2 mmol) in THF (75 mL) was added and stirred at -78°C for 2 h. It was stirred at room temperature overnight. Water (225 mL) was added, it was cooled with a water-ice bath and adjusted to pH 1 with 48% HBr. The organic phase was separated. The aqueous phase was heated to reflux for 2 h, allowed to cool and extracted with diethyl ether. The aqueous phase was taken to neutral pH with 1 N NaOH and extracted with EtOAc. The organic phase was dried over Na_2SO_4 and concentrated to dryness, to afford 28.53 g of title compound (yield: 90%).

30

¹H NMR (300 MHz, CDCl₃) δ (TMS): 4.29 (s, 2 H), 7.20 (dd, J_o = 1.6 Hz, J_m = 4.4 Hz, 2 H), 7.49 (m, 2 H), 7.58 (m, 1 H), 8.00 (d, J = 8.2 Hz, 2 H), 8.56 (dd, J_o = 1.6 Hz, J_m = 4.4 Hz, 2 H).

REFERENCE EXAMPLE 4

5 1-(4-Fluorophenyl)-2-(4-pyridyl)vinyl 4-fluorobenzoate

To a suspension of NaH (0.81 g, 18.6 mmol) in DMF (30 mL) under argon atmosphere and cooled to 0 °C, a solution of 1-(4-fluorophenyl)-2-(4-pyridyl)ethanone (2.00 g, 9.3 mmol, obtained in reference example 1) in DMF (15 mL) was added and stirred to room temperature for 30 minutes. It was then cooled to 0°C and a solution of 4-fluorobenzoyl chloride (2.95 g, 1.9 mmol) in DMF (10 mL) was added. It was stirred at room temperature overnight. Water was added and the solvent was evaporated off. The residue was dissolved in a mixture of CHCl₃ and water and the phases were separated. The aqueous phase was extracted with CHCl₃ (x3). The organic phase was washed with water (x2), dried over Na₂SO₄ and concentrated to dryness. The crude product obtained was purified by chromatography on silica gel using hexane-EtOAc mixtures of increasing polarity as eluent, to afford 0.98 g of the desired compound as a yellow solid (yield: 31%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 6.68 (s, 1 H), 7.11 (t, J = 8.6 Hz, 2 H), 7.29 (t, J = 8.6 Hz, 2 H), 7.39 (d, J = 6.0 Hz, 2 H), 7.60 (dd, J_o = 5.2 Hz, J_m = 8.8 Hz, 2 H), 8.27 (dd, J_o = 5.4 Hz, J_m = 8.8 Hz, 2 H), 8.58 (d, J = 6.0 Hz, 2 H)

REFERENCE EXAMPLE 5

1-Phenyl-2-(4-pyridyl)vinyl benzoate

Following a similar procedure to that described in reference example 4, but using 1-phenyl-2-(4-pyridyl)ethanone (obtained in reference example 3) instead of 1-(4-fluorophenyl)-2-(4-pyridyl)ethanone and benzoyl chloride instead of 4-fluorobenzoyl chloride, the title compound was obtained (yield: 62%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 6.72 (s, 1 H), 7.38 – 7.42 (complex signal, 5 H), 7.60 – 7.63 (complex signal, 4 H), 7.71 (t, J = 7.4, 1 H), 8.23 (d, J = 7.1 Hz, 2 H), 8.51 (dd, J_o = 1.5 Hz, J_m = 4.6 Hz, 2 H).

REFERENCE EXAMPLE 6

2-[2-(Methylsulfanyl)pyrimidin-4-yl]-1-[3-(trifluoromethyl)phenyl]ethanone

a) 4-Methyl-2-(methylsulfanyl)pyrimidine

To a solution of NaOH (7.46 g, 186.4 mmol) in water (120 mL), 4-methylpyrimidine-2-thiol hydrochloride (13.78 g, 84.7 mmol) was added and subsequently iodomethane (13.23 g, 93.2 mmol) was added dropwise under argon atmosphere. It was stirred at room temperature for 2 h. It was extracted with
5 CH₂Cl₂ (x2). The organic phase was dried over Na₂SO₄ and concentrated to dryness. The crude product obtained was purified by chromatography on silica gel using hexane-EtOAc mixtures of increasing polarity as eluent, to afford 10.26 g of the desired compound (yield: 88%).

b) Title compound

10 Following a similar procedure to that described in reference example 1 section b, but using *N*-methoxy-*N*-methyl-3-(trifluoromethyl)benzamide (obtained in section a of the reference example 2) instead of 4-fluoro-*N*-methoxy-*N*-methylbenzamide and 4-methyl-2-(methylsulfanyl)pyrimidine (obtained in section a of this example) instead of 4-methylpyridine, the title compound as a crude product was obtained
15 that was directly used in the following reactions (yield: quantitative).

REFERENCE EXAMPLE 7

3-(Dimethylamino)-1-(4-fluorophenyl)-2-(4-pyridyl)prop-2-en-1-one

To a solution of 1-(4-fluorophenyl)-2-(4-pyridyl)ethanone (0.30 g, 1.4 mmol, obtained in reference example 1) in anhydrous THF (5 mL), dimethyl
20 dimethylformamide acetal (0.27 g, 3.2 mmol) was added under argon atmosphere. This was stirred overnight at room temperature. The solvent was evaporated to afford 0.39 g of the title compound (yield: quantitative).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 2.79 (s, 6 H), 6.97 (t, J = 8.7 Hz, 2 H), 7.05 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H), 7.38 (s, 1 H), 7.45 (m, 2 H), 8.48 (dd, J_o = 1.5
25 Hz, J_m = 4.5 Hz, 2 H).

REFERENCE EXAMPLE 8

3-(Dimethylamino)-2-[2-(methylsulfanyl)pyrimidin-4-yl]-1-[3-(trifluoromethyl)phenyl]prop-2-en-1-one

Following a similar procedure to that described in reference example 7, but using
30 2-[2-(methylsulfanyl)pyrimidin-4-yl]-1-[3-(trifluoromethyl)phenyl]ethanone (obtained in reference example 6) instead of 1-(4-fluorophenyl)-2-(4-pyridyl)ethanone, the desired compound was obtained in the form of a crude product that was directly used in the following reactions.

REFERENCE EXAMPLE 9

1-(6-Chloropyridin-3-yl)-2-(4-pyridyl)ethanone

a) 6-Chloronicotinoyl chloride hydrochloride

A solution of 6-chloronicotinic acid (10.00 g, 63.5 mmol) in SOCl_2 (37 mL) was heated to reflux for 2 h. The SOCl_2 was evaporated to dryness, to afford 12.56 g of the desired product (yield: 93%).

b) 6-Chloro-N-methoxy-N-methylnicotinamide

Following a similar procedure to that described in reference example 1 section a, but using 6-chloronicotinoyl chloride hydrochloride (obtained in section a of this example) instead of 4-fluorobenzoyl chloride, the desired compound was obtained (yield: 71%).

c) Title compound

Following a similar procedure to that described in reference example 1 section b, but using 6-chloro-N-methoxy-N-methylnicotinamide (obtained in section b of this example) instead of 4-fluoro-N-methoxy-N-methylbenzamide, the title compound was obtained (yield: quantitative).

^1H NMR (300 MHz, CDCl_3) δ (TMS): 4.27 (s, 2 H), 7.18 (dd, $J_o = 1.5$ Hz, $J_m = 4.5$ Hz, 2 H), 7.45 (dd, $J_o = 0.6$ Hz, $J_m = 8.4$ Hz, 1 H), 8.20 (dd, $J_o = 2.5$ Hz, $J_m = 8.3$ Hz, 2 H), 8.56 (dd, $J_o = 1.6$ Hz, $J_m = 4.4$ Hz, 2 H), 8.98 (d, $J = 2.4$ Hz, 1 H).

REFERENCE EXAMPLE 10

2-(4-Fluorophenyl)-6-hydroxy-3,4'-bipyridine-5-carbonitrile

To a solution of 3-(dimethylamino)-1-(4-fluorophenyl)-2-(4-pyridyl)prop-2-en-1-one (12.77 g, 47.2 mmol, obtained in reference example 7) in DMF (175 mL), 2-cyanoacetamide (4.41 g, 52.0 mmol) was added under argon atmosphere. Then, sodium methoxide (5.35 g, 99.2 mmol) was added and heated to reflux for 1 hour. The mixture was allowed to cool, concentrated and diluted with water. The pH was adjusted to 4 with 1 N HCl. A precipitate was obtained that was filtered and dried to give 6.57 g of the desired compound as a solid (yield: 48%).

^1H NMR (300 MHz, $\text{CDCl}_3 + \text{CD}_3\text{OD}$) δ (TMS): 4.20 (s, OH + NH + CD_3OD), 6.96 (dd, $J_o = 1.6$ Hz, $J_m = 4.5$ Hz, 2 H), 7.05 (t, $J = 8.7$ Hz, 2 H), 7.23 (m, 2 H), 7.96 (s, 1 H), 8.39 (dd, $J_o = 1.4$ Hz, $J_m = 4.6$ Hz, 2 H).

REFERENCE EXAMPLE 11

2-Hydroxy-5-[2-(methylsulfanyl)pyrimidin-4-yl]-6-[3-(trifluoromethyl)phenyl]pyridine-3-carbonitrile

Following a similar procedure to that described in reference example 10, but using 3-(dimethylamino)-2-[2-(methylsulfanyl)pyrimidin-4-yl]-1-[3-(trifluoromethyl)phenyl]prop-2-en-1-one (obtained in reference example 8) instead of 3-(dimethylamino)-1-(4-fluorophenyl)-2-(4-pyridyl)prop-2-en-1-one, the title compound was obtained (yield: 20%).

¹H NMR (300 MHz, CDCl₃ + CD₃OD) δ (TMS): 3.28 (s, 3 H), 3.84 (s, OH + CD₃OD), 6.38 (d, J = 5.4 Hz, 1 H), 7.42 (d, J = 7.8 Hz, 1 H), 7.53 (t, J = 7.8 Hz, 1 H), 7.61 (s, 1 H), 7.73 (d, J = 7.8 Hz, 1 H), 8.18 (d, J = 5.1 Hz, 1 H), 8.35 (s, 1 H).

REFERENCE EXAMPLE 12

6-Chloro-2-(4-fluorophenyl)-3,4'-bipyridine-5-carbonitrile

A mixture of 2-(4-fluorophenyl)-6-hydroxy-3,4'-bipyridine-5-carbonitrile (6.57 g, 22.5 mmol, obtained in reference example 10), POCl₃ (26.3 mL, 287.5 mmol) and DMF (0.37 mL) was heated to reflux under argon atmosphere for 2 h. It was cooled with an ice bath and basified by adding 30% aqueous NH₃. The precipitate obtained was filtered and washed with water. The product was purified by chromatography on silica gel using hexane-EtOAc mixtures of increasing polarity as eluent, to afford 3.86 g of the desired product as a yellow solid (yield: 55%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 7.00 (t, J = 8.6 Hz, 2 H), 7.11 (dd, J_o = 1.6 Hz, J_m = 4.4 Hz, 2 H), 7.37 (m, 2 H), 7.98 (s, 1 H), 8.62 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H).

REFERENCE EXAMPLE 13

1-(6-Methylpyridin-3-yl)-2-(4-pyridyl) ethanone

a) N-Methoxy-6,N-dimethylnicotinamide

To a solution of 6-methylnicotinic acid (5.00 g, 36.5 mmol) in DMF (150 mL), 1-hydroxybenzotriazole (4.92 g, 36.5 mmol), N-ethyl-N-(3-dimethylaminopropyl) carbodiimide hydrochloride (8.38 g, 45.7 mmol) and 4-methylmorpholine (16.0 mL, 145.8 mmol) were added under argon atmosphere. The mixture was stirred at room temperature for 30 minutes after which N,O-dimethylhydroxylamine hydrochloride was added (3.55 g, 36.5 mmol). The reaction mixture was stirred at room temperature overnight. The solvent was evaporated off. The residue was dissolved in a mixture of CHCl₃ and 0.2 N NaHCO₃. The phases were separated

and the aqueous phase was extracted with CHCl_3 . The organic phase was dried over Na_2SO_4 and concentrated to dryness. The product was purified by chromatography on silica gel using hexane-EtOAc mixtures of increasing polarity as eluent, to afford 1.11 g of the desired product (yield: 29%).

5 b) Title compound

Following a similar procedure to that described in reference example 1 section b, but using *N*-methoxy-6,*N*-dimethylnicotinamide (obtained in section a of this example) instead of 4-fluoro-*N*-methoxy-*N*-methylbenzamide, the title compound was obtained (yield: 87%).

10 ^1H NMR (300 MHz, CDCl_3) δ (TMS): 2.64 (s, 3 H), 4.28 (s, 2 H), 7.20 (dd, $J_o = 1.4$ Hz, $J_m = 4.6$ Hz, 2 H), 7.28 (d, $J = 8.1$ Hz, 1 H), 8.15 (dd, $J_o = 2.4$ Hz, $J_m = 8.1$ Hz, 1 H), 8.57 (dd, $J_o = 1.8$ Hz, $J_m = 4.5$ Hz, 2 H), 9.10 (d, $J = 2.4$ Hz, 1 H).

REFERENCE EXAMPLE 14

2-Chloro-5-[2-(methylsulfanyl)pyrimidin-4-yl]-6-[3-(trifluoromethyl)phenyl]pyridine-3-carbonitrile

15

Following a similar procedure to that described in reference example 12 but using 2-hydroxy-5-[2-(methylsulfanyl)pyrimidin-4-yl]-6-[3-(trifluoromethyl)phenyl]pyridine-3-carbonitrile (obtained in reference example 11) instead of 2-(4-fluorophenyl)-6-hydroxy-3,4'-bipyridine-5-carbonitrile, the title compound was obtained (yield: 44%).

20

^1H NMR (300 MHz, CDCl_3) δ (TMS): 2.48 (s, 3 H), 6.62 (d, $J = 5.1$ Hz, 1 H), 7.51 (m, 2 H), 7.71 (d, $J = 7.2$ Hz, 1 H), 7.82 (s, 1 H), 8.39 (d, $J = 5.1$ Hz, 1 H), 8.42 (s, 1 H), 8.42 (s, 1 H).

REFERENCE EXAMPLE 15

25

3-Amino-5-(1-benzylpiperidin-4-yl)-2H-pyrazole

a) Methyl 1-benzylpiperidine-4-carboxylate

To a solution of methyl piperidine-4-carboxylate (10.00 g, 6.4 mmol) and triethylamine (10.32 g, 10.2 mmol) in CHCl_3 (100 mL), benzyl bromide (14.69 g, 8.6 mmol) was added under argon atmosphere while cooling with a water and ice bath. The mixture was stirred at room temperature overnight. CHCl_3 and water were added and the two phases were separated. The aqueous phase was extracted with CHCl_3 . The organic phase was dried over Na_2SO_4 and

30

concentrated to dryness, to afford 13.80 g of the desired compound as an orange solid (yield: 88%).

b) 3-(1-Benzylpiperidin-4-yl)-3-oxopropanonitrile

To a solution of BuLi (12.4 mL of a 1.6 M solution in hexane, 19.8 mmol) in THF (25 mL) cooled to -78 °C, acetonitrile (1 mL) was added dropwise under argon atmosphere. After stirring the resulting suspension for 5 min at -78 °C, a solution of methyl 1-benzylpiperidine-4-carboxylate (2.0 g, 8.1 mmol, obtained in the previous section) in THF (5 mL) was added dropwise and stirred for 30 min at -78 °C. It was allowed to reach room temperature and stirred at this temperature overnight. 1 N HCl was added to adjust the pH to 7 and the aqueous phase was extracted with CHCl₃. The organic phase was dried over Na₂SO₄ and concentrated to dryness, to afford 1.92 g of the desired compound in a solid orange form (yield: 98%).

c) Title compound

To a solution of 3-(1-benzylpiperidin-4-yl)-3-oxopropanonitrile (1.85 g, 7.6 mmol, obtained in the previous section) in EtOH (77 mL), hydrazine monohydrate (0.74 mL, 15.3 mmol) was added under argon atmosphere. The mixture was heated to reflux overnight. The solvent was evaporated and the residue was dissolved in a mixture of water-CHCl₃. The aqueous phase was extracted with CHCl₃. The organic phase was dried over Na₂SO₄ and concentrated to dryness. The crude product obtained was purified by chromatography on silica gel using hexane-EtOAc mixtures of increasing polarity as eluent, to afford 0.46 g of the desired compound (yield: 23%)

¹H NMR (300 MHz, CD₃OD) δ (TMS): 1.72 (m, 2 H), 1.90 (m, 2 H), 2.15 (m, 2 H), 2.56 (m, 1 H), 2.97 (m, 2 H), 3.57 (s, 2 H), 4.89 (broad s, NH + NH₂ + H₂O), 5.43 (s, 1 H), 7.27 – 7.34 (complex signal, 5 H).

REFERENCE EXAMPLE 16

3-(Dimethylamino)-1-(4-fluorophenyl)-2-(4-pyridyl)but-2-en-1-one

Following a similar procedure to that described in reference example 7, but using dimethylacetamide dimethyl acetal instead of dimethylformamide dimethyl acetal, the desired compound was obtained in the form of a crude product that was directly used in the following reactions.

¹H NMR (300 MHz, CDCl₃) δ (TMS): 2.15 (s, 3 H), 3.00 (s, 6 H), 6.80 (m, 4 H), 7.45 (m, 2 H), 8.30 (d, J = 8.0 Hz, 2 H).

REFERENCE EXAMPLE 17

2-(4-Fluorophenyl)-6-hydroxy-4-methyl-3,4'-bipyridine-5-carbonitrile

5 Following a similar procedure to that described in reference example 10, but using 3-(dimethylamino)-1-(4-fluorophenyl)-2-(4-pyridyl)but-2-en-1-one (obtained in reference example 16) instead of 3-dimethylamino-1-(4-fluorophenyl)-2-(4-pyridyl)prop-2-en-1-one, the title compound was obtained (yield: 21%).

10 ¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.55 (broad s, OH + H₂O), 2.30 (s, 3 H), 6.97 (m, 4 H), 7.25 (m, 2 H), 8.52 (m, 2 H).

REFERENCE EXAMPLE 18

6-Chloro-2-(4-fluorophenyl)-4-methyl-3,4'-bipyridine-5-carbonitrile

Following a similar procedure to that described in reference example 12, but using 2-(4-fluorophenyl)-6-hydroxy-4-methyl-3,4'-bipyridine-5-carbonitrile (obtained in reference example 17) instead of 2-(4-fluorophenyl)-6-hydroxy-3,4'-bipyridine-5-carbonitrile, the title compound was obtained (yield: 52%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 2.41 (s, 3 H), 6.90 (m, 2 H), 7.09 (d, J = 9.0 Hz, 2 H), 7.25 (m, 2 H), 8.62 (m, J = 4.0 Hz, 2 H).

REFERENCE EXAMPLE 19

1-(4-Fluorophenyl)-2-[2-(methylsulfonyl)pyrimidin-4-yl]ethanone

20 To a solution of 4-methyl-2-(methylsulfonyl)pyrimidine (21.00 g, 150.0 mmol, obtained in reference example 6, section a) and ethyl 4-fluorobenzoate (25.14 g, 150.0 mmol) in THF (300 mL) under argon atmosphere, a solution of sodium bis(trimethylsilyl)amide (150 mL of a 2 M solution in THF, 300 mmol) in THF (150 mL) was added dropwise while cooling with an ice-bath. It was stirred at room temperature for 2 h. Saturated NH₄Cl was added and the solvent was evaporated. The residue was taken up in a mixture of EtOAc and water and the phases were separated. The aqueous phase was reextracted with EtOAc. The combined organic phases were washed with brine, dried over Na₂SO₄ and concentrated to dryness, to afford 36.36 g of the title compound (yield: 93%).

30 ¹H NMR (300 MHz, CDCl₃) δ (TMS): 2.52 (ketone: s, 3 H), 2.61 (enol: s, 3 H), 4.35 (ketone: s, 2 H), 5.92 (enol: s, 1 H), 6.64 (enol: d, J = 5.7 Hz, 1 H), 6.95 (ketone: d,

J = 5.1 Hz, 1 H), 7.08.- 7.19 (m, 2H), 7.83 (enol: m, 2 H), 8.07 (ketone: m, 2 H), 8.31 (enol: d, J = 5.7 Hz, 1 H), 8.56 (ketone: d, J = 5.1 Hz, 1 H).

REFERENCE EXAMPLE 20

3-(Dimethylamino)-1-(4-fluorophenyl)-2-[2-(methylsulfanyl)pyrimidin-4-yl]prop-2-en-1-one

5

Following a similar procedure to that described in reference example 7, but using 1-(4-fluorophenyl)-2-[2-(methylsulfanyl)pyrimidin-4-yl]ethanone (obtained in reference example 19) instead of 1-(4-fluorophenyl)-2-(4-pyridyl)ethanone, the title compound was obtained.

10 ¹H NMR (300 MHz, CDCl₃) δ (TMS): 2.50 (s, 3 H), 2.96 (s, 6 H), 6.20 - 8.20 (complex signal, 7 H).

REFERENCE EXAMPLE 21

6-(4-Fluorophenyl)-2-(hydroxy)-5-(2-methylsulfanylpyrimidin-4-yl)pyridine-3-carbonitrile

15 Following a similar procedure to that described in reference example 10, but using 3-(dimethylamino)-1-(4-fluorophenyl)-2-[2-(methylsulfanyl)pyrimidin-4-yl]prop-2-en-1-one (obtained in reference example 20) instead of 3-(dimethylamino)-1-(4-fluorophenyl)-2-(4-pyridyl)prop-2-en-1-one, the title compound was obtained (yield: 91%).

20 LC-MS (method 1): t_R = 7.09 min; m/z = 338.9 [M+H]⁺

REFERENCE EXAMPLE 22

2-Chloro-6-(4-fluorophenyl)-5-(2-methylsulfanylpyrimidin-4-yl)pyridine-3-carbonitrile

25 A mixture of 6-(4-fluorophenyl)-2-(hydroxy)-5-(2-methylsulfanylpyrimidin-4-yl)pyridine-3-carbonitrile (48.84 g, 144.6 mmol, obtained in reference example 21), POCl₃ (166 mL, 1.8 mol) and DMF (2.2 mL) was heated to 100 °C for 2 h. It was allowed to cool to room temperature and concentrated. It was cooled with an acetone-CO₂ bath and EtOAc and ice were subsequently added. The organic phase was decanted, washed with saturated NaHCO₃, dried over Na₂SO₄ and concentrated to dryness. The crude obtained was purified by chromatography on silica gel using hexane-EtOAc mixtures of increasing polarity, to afford 31.00 g of the title compound (yield: 60%).

30

¹H NMR (300 MHz, CDCl₃) δ (TMS): 2.54 (s, 3 H), 6.60 (d, J = 5.1 Hz, 1 H), 7.08 (t, J = 8.5 Hz, 2 H), 7.44 (m, 2 H), 8.37 (d, J = 5.1 Hz, 1 H), 8.39 (s, 1 H).

REFERENCE EXAMPLE 23

2-(2-Chloropyridin-4-yl)-1-(4-fluorophenyl)ethanone

- 5 Following a similar procedure to that described in reference example 19, but starting from 2-chloro-4-methylpyridine and ethyl 4-fluorobenzoate, the title compound was obtained.

LC-MS (method 1): t_R = 7.96 min; m/z = 250.0, 252.0 [M+H]⁺

REFERENCE EXAMPLE 24

- 10 2-[2-(Methylsulfanyl)pyrimidin-4-yl]-1-phenylethanone

Following a similar procedure to that described in reference example 19, but starting from 4-methyl-2-(methylsulfanyl)pyrimidine (obtained in reference example 6, section a) and ethyl benzoate, the title compound was obtained.

- 15 ¹H NMR (300 MHz, CDCl₃) δ (TMS): 2.52 (ketone: s, 3 H), 2.62 (enol: s, 3 H), 4.39 (ketone: s, 2 H), 5.99 (enol: s, 1 H), 6.65 (enol: d, J = 5.7 Hz, 1 H), 6.98 (ketone: d, J = 5.1 Hz, 1 H), 7.40 - 7.51 (m, 3 H), 7.85 (enol: m, 2 H), 8.03 (ketone: m, 2 H), 8.32 (enol: d, J = 5.7 Hz, 1 H), 8.46 (ketone: d, J = 5.1 Hz, 1 H).

REFERENCE EXAMPLE 25

- 20 3-(Dimethylamino)-2-[2-(methylsulfanyl)pyrimidin-4-yl]-1-phenylprop-2-en-1-one

Following a similar procedure to that described in reference example 7, but starting from 2-[2-(methylsulfanyl)pyrimidin-4-yl]-1-phenylethanone (obtained in reference example 24), the title compound was obtained.

- 25 ¹H NMR (300 MHz, CDCl₃) δ (TMS): 2.50 (s, 3 H), 2.90 (s, 6 H), 6.20 - 8.00 (complex signal, 8 H).

REFERENCE EXAMPLE 26

2-Hydroxy-5-(2-methylsulfanylpyrimidin-4-yl)-6-phenylpyridine-3-carbonitrile

- 30 Following a similar procedure to that described in reference example 10, but using 3-(dimethylamino)-2-[2-(methylsulfanyl)pyrimidin-4-yl]-1-phenylprop-2-en-1-one (obtained in reference example 25) instead of 3-dimethylamino-1-(4-fluorophenyl)-2-(4-pyridyl)prop-2-en-1-one, the title compound was obtained.

LC-MS (method 1): t_R = 6.90 min; m/z = 320.9 [M+H]⁺

REFERENCE EXAMPLE 27

1-[6-(4-Fluorophenyl)-2-hydroxy-5-(2-methylsulfanylpirimidin-4-yl)pyridin-3-yl]ethanone

Following a similar procedure to that described in reference example 10, but starting from 3-(dimethylamino)-1-(4-fluorophenyl)-2-[2-(methylsulfanyl)pyrimidin-4-yl]prop-2-en-1-one (obtained in reference example 20) and 3-oxobutylamide, the title compound was obtained.

¹H NMR (300 MHz, CDCl₃) δ (TMS): 2.43 (s, 3 H), 2.66 (enol: s, 3 H), 6.53 (d, J = 5.1 Hz, 1 H), 7.12 (t, J = 8.4 Hz, 2 H), 7.42 (m, 2 H), 8.29 (d, J = 5.1 Hz, 1 H), 8.63 (s, 1 H).

10

REFERENCE EXAMPLES 28-29

Following a similar procedure to that described in reference example 12, but starting from the appropriate compounds in each case, the compounds in the following table were obtained:

Reference example	Compound name	Starting compounds	LC-MS		
			Method	t _R (min)	m/z [M+H] ⁺
28	2-Chloro-5-(2-methylsulfanylpirimidin-4-yl) 6-phenylpyridine-3-carbonitrile	Reference example 26	1	10.18	338.9
29	4-[6-Chloro-5-(1-chlorovinyl)-2-(4-fluorophenyl)pyridin-3-yl]-2-methylsulfanylpirimidine	Reference example 27	1	11.62	391.9 393.9 395.9

15

REFERENCE EXAMPLE 30

1-(4-Fluorophenyl)-2-pyrimidin-4-ylethanone

To a suspension of NaH (2.26 g 50%, 47.7 mmol) in DMF (92 mL) under argon atmosphere and cooled to 0 °C, 4-methylpyrimidine (3.00 g, 31.9 mmol) was added slowly. Then, ethyl 4-fluorobenzoate (6.40 g, 38.2 mmol) was added and it was stirred at room temperature overnight. Water was added and the solvent was evaporated. The residue was taken up in a mixture of EtOAc and brine. The phases were separated and the aqueous phase was reextracted with EtOAc. The combined organic phases were dried over Na₂SO₄ and concentrated to dryness. The crude product obtained by chromatography on silica gel using hexane-EtOAc

20

mixtures of increasing polarity as eluent, to afford 3.30 g of the desired compound (yield: 48%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 4.11 (ketone: s, 2 H), 5.94 (enol: s, 1 H), 6.94 (enol: d, J = 5.4 Hz, 1 H), 7.08 - 7.16 (m, 2 H), 7.37 (ketone: d, J = 5.1 Hz, 1 H),
5 7.89 (enol: m, 2 H), 8.08 (ketone: m, 2 H), 8.42 (enol: d, J = 5.4 Hz, 1 H), 8.69 (ketone: d, J = 5.1 Hz, 1 H), 8.81 (enol: s, 1 H), 9.17 (ketone: s, 1 H).

EXAMPLE 1

4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)-1H-pyrazolo[3,4-b]pyridine

METHOD A

10 In a volumetric flask, 1-(4-fluorophenyl)-2-(4-pyridyl)ethanone (23.56 g, 109.4 mmol, obtained in reference example 1) and 2-methoxyethanol (150 mL) were introduced. A solution of 3-amino-2H-pyrazole (10.00 g, 120.3 mmol) in 2-methoxyethanol (170 mL) and 37% HCl (3.23 g, 32.8 mmol) were added under argon atmosphere. This was heated to reflux for 3 days. It was allowed to cool and
15 concentrated. The solid obtained was dissolved in CHCl₃ (400 mL) and MeOH (50 mL) and washed with 0.1 N HCl (300 mL) and 1 N NaOH (300 mL). The organic phase was dried over Na₂SO₄ and concentrated to dryness, to afford 9.93 g of the desired product in solid cream form (yield: 47%)

METHOD B

20 To a solution of 3-amino-2H-pyrazole (60 mg, 71.8 mmol) in EtOH (2 mL) and 1 drop of 37% HCl, 1-(4-fluorophenyl)-2-(4-pyridyl)vinyl 4-fluorobenzoate (0.22 g, 65.0 mmol, obtained in reference example 4) was added under argon atmosphere. This was heated to reflux for 3 days. The mixture was diluted with CHCl₃ and MeOH. It was washed with saturated NaHCO₃. The aqueous phase was extracted
25 with CHCl₃ (x2). The organic phase was dried over Na₂SO₄ and concentrated to dryness. The crude product obtained was purified by chromatography on silica gel using hexane-EtOAc mixtures of increasing polarity as eluent, to afford 58 mg of the desired product in a solid white form (yield: 23%).

¹H NMR (300 MHz, CDCl₃ + CD₃OD) δ (TMS): 4.08 (s, NH + CD₃OD), 6.80 - 7.01 (complex signal, 6 H), 7.21 (m, 2 H), 7.28 (m, 2 H), 7.95 (s, 1 H), 8.27 (dd, J_o = 1.4 Hz, J_m = 4.6 Hz, 2 H).

EXAMPLE 2

4,6-Diphenyl-5-(4-pyridyl)-1H-pyrazolo[3,4-b]pyridine

Following a similar procedure to that described in example 1 method B, but using 1-phenyl-2-(4-pyridyl)vinyl benzoate (obtained in reference example 5) instead of 1-(4-fluorophenyl)-2-(4-pyridyl)vinyl 4-fluorobenzoate, the title compound was obtained in solid white form (yield: 37%).

- 5 ¹H NMR (300 MHz, CDCl₃) δ (TMS): 4.08 (s, NH + H₂O), 6.85 (d, J = 6.0 Hz, 2 H), 7.12 - 7.31 (complex signal, 10 H), 7.98 (s, 1 H), 8.29 (d, J = 5.8 Hz, 2 H).

EXAMPLE 3

5-(4-Pyridyl)-4,6-bis[3-(trifluoromethyl)phenyl]-1H-pyrazolo[3,4-b]pyridine

- Following a similar procedure to that described in example 1 method A, but using
10 2-(4-pyridyl)-1-[3-(trifluoromethyl)phenyl]ethanone (obtained in reference example 2) instead of 1-(4-fluorophenyl)-2-(4-pyridyl)ethanone, the title compound was obtained (yield: 10%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.57 (s, NH + H₂O), 6.86 (broad s, 2 H), 7.30 - 7.60 (complex signal, 8 H), 7.99 (s, 1 H), 8.35 (broad s, 2 H).

15

EXAMPLE 4

4,6-Bis(4-fluorophenyl)-3-methyl-5-(4-pyridyl)-1H-pyrazolo[3,4-b]pyridine

Following a similar procedure to that described in example 1 method B, but using 3-amino-5-methyl-2H-pyrazole instead of 3-amino-2H-pyrazole, the title compound was obtained in solid white form (yield: 19%).

- 20 ¹H NMR (300 MHz, CDCl₃ + CD₃OD) δ (TMS): 2.03 (broad s, 3 H), 4.08 (s, NH + CD₃OD), 6.81 (m, 2 H), 6.96 (m, 2 H), 7.01 (m, 2 H), 7.04 (m, 2 H), 7.29 (m, 2 H), 8.23 (m, 2 H).

EXAMPLE 5

4,6-Diphenyl-3-methyl-5-(4-pyridyl)-1H-pyrazolo[3,4-b]pyridine

- 25 Following a similar procedure to that described in example 1 method B, but using 3-amino-5-methyl-2H-pyrazole instead of 3-amino-2H-pyrazole and 1-phenyl-2-(4-pyridyl)vinyl benzoate (obtained in reference example 5) instead of 1-(4-fluorophenyl)-2-(4-pyridyl)vinyl 4-fluorobenzoate, the title compound was obtained in solid white form (yield: 16%).

- 30 ¹H NMR (300 MHz, CDCl₃) δ (TMS): 2.02 (s, 3 H), 2.02 (s, NH + H₂O), 6.83 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H), 7.13 (m, 2 H), 7.23 - 7.33 (complex signal, 6 H), 8.25 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H).

EXAMPLE 6

2-Ethyl-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine**EXAMPLE 7****1-Ethyl-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine**

A suspension of 4,6-bis(4-fluorophenyl)-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine (0.39 g, 0.8 mmol, obtained in example 1), KOH (0.05 g, 0.8 mmol) and crown ether 18-C-6 (0.01 g, 0.03 mmol) in toluene (3 mL) was heated to 100 °C for 2 h. A solution of iodoethane (0.18 g, 1.2 mmol) in toluene (1 mL) was added and stirred at 100 °C for 2 days. It was allowed to cool, water and EtOAc were added and the phases were separated. The aqueous phase was extracted with EtOAc. The organic phase was dried over Na₂SO₄ and concentrated to dryness. The crude product obtained was purified by chromatography on silica gel using hexane-EtOAc mixtures of increasing polarity as eluent, to afford 0.20 g of 2-ethyl-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine (yield: 61%) and 28 mg of 1-ethyl-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine (yield: 9%).

Example 6: ¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.68 (t, *J* = 7.4 Hz, 3 H), 4.51 (c, *J* = 7.2 Hz, 2 H), 6.82 (dd, *J*_o = 1.6 Hz, *J*_m = 4.4 Hz, 2 H), 6.89 (t, *J* = 8.7 Hz, 2 H), 7.00 (t, *J* = 8.7 Hz, 2 H), 7.15 (m, 2 H), 7.30 (m, 2 H), 7.78 (s, 1H), 8.31 (dd, *J*_o = 1.6 Hz, *J*_m = 4.5 Hz, 2 H).

Example 7: ¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.61 (t, *J* = 7.4 Hz, 3 H), 4.67 (c, *J* = 7.2 Hz, 2 H), 6.81 (dd, *J*_o = 1.6 Hz, *J*_m = 4.4 Hz, 2 H), 6.96 (t, *J* = 8.7 Hz, 2 H), 7.01 (t, *J* = 8.7 Hz, 2 H), 7.13 (m, 2 H), 7.29 (m, 2 H), 7.86 (s, 1H), 8.32 (dd, *J*_o = 1.5 Hz, *J*_m = 4.5 Hz, 2 H).

EXAMPLE 8**4,6-Bis(4-fluorophenyl)-2-methyl-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine**

25

EXAMPLE 9**4,6-Bis(4-fluorophenyl)-1-methyl-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine**

Following a similar procedure to that described in example 6 and 7, but using iodomethane instead of iodoethane, the title compounds were obtained.

Example 8: yield: 52%; ¹H NMR (300 MHz, CDCl₃) δ (TMS): 4.25 (s, 3 H), 6.81 (d, *J* = 5.3 Hz, 2 H), 6.95 (t, *J* = 8.6 Hz, 2 H), 6.98 (t, *J* = 8.5 Hz, 2 H), 7.12 (m, 2 H), 7.28 (m, 2 H), 7.77 (s, 1H), 8.29 (d, *J* = 5.2 Hz, 2 H).

30

Example 9: yield: 5%; ¹H NMR (300 MHz, CDCl₃) δ (TMS): 4.20 (s, 3 H), 6.81 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H), 6.90 (t, J = 8.7 Hz, 2 H), 6.98 (t, J = 8.6 Hz, 2 H), 7.12 (m, 2 H), 7.26 (m, 2 H), 7.84 (s, 1H), 8.29 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H).

EXAMPLE 10

5 **4,6-Bis(4-fluorophenyl)-2,3-dimethyl-5-(4-pyridyl)pyrazolo[3,4-b]pyridine**

EXAMPLE 11

4,6-Bis(4-fluorophenyl)-1,3-dimethyl-5-(4-pyridyl)pyrazolo[3,4-b]pyridine

Following a similar procedure to that described in examples 6 and 7, but using 4,6-bis(4-fluorophenyl)-3-methyl-5-(4-pyridyl)-1H-pyrazolo[3,4-b]pyridine (obtained in
10 example 4) instead of 4,6-bis(4-fluorophenyl)-5-(4-pyridyl)-1H-pyrazolo[3,4-b]pyridine and iodomethane instead of iodoethane, the title compounds were obtained.

Example 10: yield: 53%; ¹H NMR (300 MHz, CDCl₃) δ (TMS): 2.03 (s, 3 H), 4.13 (s, 3 H), 6.79 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H), 6.88 (t, J = 8.8 Hz, 2 H), 6.98 –
15 7.11 (complex signal, 4 H), 7.30 (m, 2 H), 8.26 (dd, J_o = 1.6 Hz, J_m = 4.4 Hz, 2 H).

Example 11: yield: 30%; ¹H NMR (300 MHz, CDCl₃ + CD₃OD) δ (TMS): 2.01 (broad s, 3 H), 4.15 (broad s, 3 H), 6.81 (m, 2 H), 6.93 (m, 2 H), 7.00 (m, 2 H), 7.07 (m, 2 H), 7.30 (m, 2 H), 8.23 (m, 2 H).

EXAMPLE 12

20 **2-[2-[1-(tert-Butoxycarbonyl)piperidin-4-yl]ethyl]-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridine**

EXAMPLE 13

1-[2-[1-(tert-Butoxycarbonyl)piperidin-4-yl]ethyl]-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridine

25 **a) 2-[1-(tert-Butoxycarbonyl)piperidin-4-yl]ethanol**

To a solution of 2-(4-piperidyl)ethanol (9.63 g, 74.5 mmol) in DMF (100 mL) at 0 °C, di-tert-butyl dicarbonate (16.26 g, 74.5 mol) was added slowly. The mixture was stirred overnight at room temperature. The solvent was concentrated and the residue was dissolved in a mixture of EtOAc and water. The phases were
30 separated. The organic phase was dried over Na₂SO₄ and concentrated to dryness, to afford 15.09 g of the desired product (yield: 88%).

b) 2-[1-(tert-Butoxycarbonyl)piperidin-4-yl]ethyl methanesulfonate

To a solution of 2-[1-(*tert*-butoxycarbonyl)piperidin-4-yl]ethanol (7.50 g, 32.7 mmol, obtained in section a) in CHCl_3 (180 mL), triethylamine (4.6 mL) was added under argon atmosphere and cooled to 0°C. Then, methanesulfonyl chloride (2.6 mL, 32.7 mmol) was added dropwise. The mixture was stirred overnight at room temperature. Water was added and the phases were separated. The aqueous phase was extracted with CHCl_3 (x3). The organic phase was dried over Na_2SO_4 and concentrated to dryness, to afford 11.18 g of the desired compound (yield: quantitative).

c) Title compounds

Following a similar procedure to that described in example 6 and 7, but using 2-[1-(*tert*-butoxycarbonyl)piperidin-4-yl]ethyl methanesulfonate (obtained in section b) instead of iodoethane, the title compounds were obtained.

Example 12: yield: 14%; ^1H NMR (300 MHz, CDCl_3) δ (TMS): 1.00 -1.40 (complex signal, 3 H), 1.45 (s, 9 H), 1.70 (m, 2 H), 2.66 (m, 2 H), 2.05 (m, 2 H), 4.09 (m, 2 H), 4.90 (t, $J = 7.4$ Hz, 2 H), 6.83 (dd, $J_o = 1.6$ Hz, $J_m = 4.4$ Hz, 2 H), 6.90 (t, $J = 8.7$ Hz, 2 H), 7.00 (t, $J = 8.7$ Hz, 2 H), 7.14 (m, 2 H), 7.28 (m, 2 H), 7.77 (s, 1 H), 8.31 (dd, $J_o = 1.6$ Hz, $J_m = 4.4$ Hz, 2 H).

Example 13: yield: 29%; ^1H NMR (300 MHz, CDCl_3) δ (TMS):): 1.00 -1.40 (complex signal, 3 H), 1.46 (s, 9 H), 1.82 (m, 2 H), 1.98 (m, 2 H), 2.67 (m, 2 H), 4.08 (m, 2 H), 4.65 (t, $J = 7.0$ Hz, 2 H), 6.82 (dd, $J_o = 1.6$ Hz, $J_m = 4.5$ Hz, 2 H), 6.93 (t, $J = 8.7$ Hz, 2 H), 7.01 (t, $J = 8.7$ Hz, 2 H), 7.25 (m, 2 H), 7.28 (m, 2 H), 7.86 (s, 1 H), 8.32 (dd, $J_o = 1.5$ Hz, $J_m = 4.5$ Hz, 2 H).

EXAMPLE 14

2-[1-(*tert*-Butoxycarbonyl)piperidin-4-yl]-4,6-bis(4-fluorophenyl)-3-methyl-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine

EXAMPLE 15

1-[1-(*tert*-Butoxycarbonyl)piperidin-4-yl]-4,6-bis(4-fluorophenyl)-3-methyl-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine

a) 1-(*tert*-Butoxycarbonyl)piperidin-4-yl methanesulfonate

Following a similar procedure to that described in examples 12 and 13 section b, but using (1-*tert*-butoxycarbonyl)piperidin-4-ol instead of 2-[1-(*tert*-butoxycarbonyl)piperidin-4-yl]ethanol, the desired compound was obtained (yield: 97%).

b) Title compounds

Following a similar procedure to that described in examples 6 and 7, but using 4,6-bis(4-fluorophenyl)-3-methyl-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine (obtained in example 4) instead of 4,6-bis(4-fluorophenyl)-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine and 1-(*tert*-butoxycarbonyl)piperidin-4-yl methanesulfonate (obtained in section a) instead of iodoethane, the title compounds were obtained.

Example 14: yield: 13%; ¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.51 (s, 9 H), 1.97 (m, 2 H), 2.06 (s, 3 H), 2.45 (m, 2 H), 2.94 (m, 2 H), 4.35 (m, 3 H), 6.78 (d, J = 6.0 Hz, 2 H), 6.87 (t, J = 8.7 Hz, 2 H), 7.01 (t, J = 8.6 Hz, 2 H), 7.10 (m, 2 H), 7.33 (m, 2 H), 8.26 (d, J = 6.0 Hz, 2 H).

Example 15: yield: 52%; ¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.54 (s, 9H), 2.00 (s, 3 H), 2.00 (m, 2 H), 2.31 (m, 2 H), 2.99 (m, 2 H), 4.33 (m, 2 H), 5.10 (m, 1 H), 6.78 (dd, J_o = 1.8 Hz, J_m = 4.5 Hz, 2 H), 6.92 (t, J = 8.7 Hz, 2 H), 6.99 (t, J = 8.7 Hz, 2 H), 7.08 (m, 2 H), 7.30 (m, 2 H), 8.27 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H).

EXAMPLE 16

2-(3-Chloropropyl)-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine

EXAMPLE 17

1-(3-Chloropropyl)-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine

Following a similar procedure to that described in examples 6 and 7, but using 1-bromo-3-chloropropane instead of iodoethane, the title compounds were obtained.

Example 16: yield: 28%; ¹H NMR (300 MHz, CDCl₃) δ (TMS): 2.54 (m, 2 H), 3.52 (t, J = 6.0 Hz, 2 H), 4.61 (t, J = 6.0 Hz, 2 H), 6.81 (dd, J_o = 1.6 Hz, J_m = 4.4 Hz, 2 H), 6.86 (t, J = 8.8 Hz, 2 H), 6.98 (t, J = 8.7 Hz, 2 H), 7.13 (m, 2 H), 7.28 (m, 2 H), 7.83 (s, 1 H), 8.29 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H).

Example 17: yield: 19%; ¹H NMR (300 MHz, CDCl₃) δ (TMS): 2.51 (m, 2 H), 3.61 (t, J = 6.3 Hz, 2 H), 4.78 (t, J = 6.4 Hz, 2 H), 6.82 (dd, J_o = 1.6 Hz, J_m = 4.4 Hz, 2 H), 6.92 (t, J = 8.7 Hz, 2 H), 7.01 (t, J = 8.7 Hz, 2 H), 7.14 (m, 2 H), 7.26 (m, 2 H), 7.86 (s, 1 H), 8.32 (dd, J_o = 1.6 Hz, J_m = 4.4 Hz, 2 H).

EXAMPLE 18

3-[4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]piperidin-2-yl]propan-1-ol

METHOD A

a) **4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)-2-[3-(tetrahydropyran-2-yloxy)propyl]pyrazolo[3,4-b]pyridine**

A suspension of 4,6-bis(4-fluorophenyl)-5-(4-pyridyl)-1H-pyrazolo[3,4-b]pyridine (0.30 g, 0.8 mmol, obtained in example 1), KOH (0.05 g, 0.8 mmol) and crown ether 18-C-6 (0.01 g, 0.03 mmol) in toluene (10 mL) was heated to 100 °C for 1 h. 2-(3-Bromopropoxy)tetrahydropyran (0.17 g, 0.8 mmol) was added and stirred at 100 °C for 24 h. It was allowed to cool, water and EtOAc were added and the phases were separated. The aqueous phase was extracted with EtOAc. The organic phase was dried over Na₂SO₄ and concentrated to dryness. The crude product obtained was purified by chromatography on silica gel using hexane-EtOAc mixtures of increasing polarity as eluent, to afford 0.22 g of the desired compound (yield: 54%).

LC-MS (método 1): t_R = 7.60 min; m/z = 527.2 [M+H]⁺.

b) **Title compound**

A solution of 4,6-bis(4-fluorophenyl)-5-(4-pyridyl)-2-[3-(tetrahydropyran-2-yloxy)propyl]pyrazolo[3,4-b]pyridine (0.22 g, 0.42 mmol, obtained in section a) in a mixture 4:2:1 of AcOH:THF:H₂O (9 mL) was heated to a 55 °C for 3 h. It was allowed to cool and concentrated. Saturated NaHCO₃ and 1 N NaOH were added to the residue and it was extracted with EtOAc. The organic phase was dried over Na₂SO₄ and concentrated to dryness, to afford 0.15 g of the title compound (yield: 83%).

LC-MS (método 1): t_R = 5.37 min; m/z = 443.1 [M+H]⁺.

EXAMPLE 18

3-[4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridin-2-yl]propan-1-ol

EXAMPLE 19

3-[4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridin-1-yl]propan-1-ol

METHOD B

Following a similar procedure to that described in examples 6 and 7, but using 3-iodopropanol instead of iodoethane, the title compounds were obtained.

Example 18: yield: 33%; ¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.58 (s, OH + H₂O), 2.17 (m, 2 H), 3.71 (m, 2 H), 4.63 (t, J = 6.4 Hz, 2 H), 6.83 (dd, J_o = 1.6 Hz, J_m =

4.4 Hz, 2 H), 6.90 (t, J = 8.7 Hz, 2 H), 7.00 (t, J = 8.7 Hz, 2 H), 7.15 (m, 2 H), 7.29 (m, 2 H), 7.82 (s, 1 H), 8.33 (dd, $J_o = 1.5$ Hz, $J_m = 4.5$ Hz, 2 H).

Example 19: yield: 21%; ^1H NMR (300 MHz, CDCl_3) δ (TMS): 1.57 (s, OH + H_2O), 2.17 (m, J = 5.9 Hz, 2 H), 3.58 (m, 2 H), 4.78 (t, J = 6.0 Hz, 2 H), 6.82 (dd, $J_o = 1.5$ Hz, $J_m = 4.5$ Hz, 2 H), 6.94 (t, J = 8.7 Hz, 2 H), 7.02 (t, J = 8.7 Hz, 2 H), 7.15 (m, 2 H), 7.27 (m, 2 H), 7.90 (s, 1 H), 8.34 (dd, $J_o = 1.5$ Hz, $J_m = 4.5$ Hz, 2 H).

EXAMPLE 20

2-[1-(*tert*-Butoxycarbonyl)piperidin-4-yl]-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine

EXAMPLE 21

1-[1-(*tert*-Butoxycarbonyl)piperidin-4-yl]-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine

Following an analogous procedure to that described in examples 6 and 7, but using 1-(*tert*-butoxycarbonyl)piperidin-4-yl methanesulfonate (obtained in example 14 section a) instead of iodoethane, the title compounds were obtained.

Example 20: yield: 30%; ^1H NMR (300 MHz, CDCl_3) δ (TMS): 1.48 (s, 9 H), 2.00-2.20 (m, 4 H), 2.87 (m, 2 H), 4.21 (m, 2 H), 4.50 (m, 1 H), 6.73 (dd, $J_o = 1.5$ Hz, $J_m = 4.5$ Hz, 2 H), 6.81 (t, J = 8.7 Hz, 2 H), 6.92 (t, J = 8.7 Hz, 2 H), 7.03 (m, 2 H), 7.20 (m, 2 H), 7.73 (s, 1 H), 8.23 (dd, $J_o = 1.4$ Hz, $J_m = 4.4$ Hz, 2 H).

Example 21: yield: 28%; ^1H NMR (300 MHz, CDCl_3) δ (TMS): 1.46 (s, 9 H), 2.10 – 2.30 (complex signal, 4 H), 2.96 (m, 2 H), 4.30 (m, 2 H), 4.60 (m, 1 H), 6.82 (dd, $J_o = 1.5$ Hz, $J_m = 4.5$ Hz, 2 H), 6.89 (t, J = 8.7 Hz, 2 H), 7.00 (t, J = 8.7 Hz, 2 H), 7.14 (m, 2 H), 7.31 (m, 2 H), 7.82 (s, 1 H), 8.31 (dd, $J_o = 1.5$ Hz, $J_m = 4.5$ Hz, 2 H).

EXAMPLE 22

2-Methyl-4,6-diphenyl-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine

EXAMPLE 23

1-Methyl-4,6-diphenyl-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine

To a suspension of 4,6-diphenyl-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine (0.10 g, 0.3 mmol, obtained in example 2) in acetone (1 mL), KOH (21 mg, 0.4 mmol) was added under argon atmosphere. Then, a solution of iodomethane (47 mg, 0.3 mmol) in acetone (0.1 mL) was added and stirred overnight at room temperature. Water was added and extracted with CHCl_3 . The organic phase was dried over Na_2SO_4 and concentrated to dryness. The crude product obtained was purified by

chromatography on silica gel using hexane-EtOAc mixtures of increasing polarity as eluent, to afford 47 mg of 2-methyl-4,6-diphenyl-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine (yield: 47%) and 38 mg of 1-methyl-4,6-diphenyl-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine (yield: 38%).

5 Example 22: ¹H NMR (300 MHz, CDCl₃) δ (TMS): 4.24 (s, 3 H), 6.83 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H), 7.16 – 7.21 (complex signal, 4 H), 7.26 – 7.31 (complex signal, 6 H), 7.77 (s, 1H), 8.24 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H).

Example 23: ¹H NMR (300 MHz, CDCl₃) δ (TMS): 4.23 (s, 3 H), 6.81 (dd, J_o = 1.6 Hz, J_m = 4.4 Hz, 2 H), 7.17 (m, 2 H), 7.22 – 7.32 (complex signal, 8 H), 7.89 (s, 1H), 8.26 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H).

EXAMPLE 24

4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)-2-[2-(tetrahydropyran-2-yloxy)ethyl]pyrazolo[3,4-*b*]pyridine

Following a similar procedure to that described in examples 6 and 7, but using 2-
15 (2-bromoethoxy)tetrahydropyran instead of iodoethane, the title compound was obtained (yield: 50%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.48 – 1.63 (complex signal, 6 H), 3.47 (m, 1 H), 3.67 (m, 1 H), 4.02 (m, 1 H), 4.22 (m, 1 H), 4.57 (m, 1 H), 4.65 (m, 2 H), 6.83 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H), 6.90 (t, J = 8.7 Hz, 2 H), 7.00 (t, J = 8.6 Hz, 2 H), 7.14 (m, 2 H), 7.31 (m, 2 H), 7.92 (s, 1 H), 8.32 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H).

EXAMPLE 25

2-[4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-1-yl]ethanol

Following a similar procedure to that described in examples 22 and 23, but using
25 4,6-bis(4-fluorophenyl)-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine (obtained in example 1) instead of 4,6-diphenyl-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine and 2-bromoethanol instead of iodomethane, the title compound was obtained.

¹H NMR (300 MHz, CDCl₃) δ (TMS): 4.19 (m, 2 H), 4.32 (m, OH), 4.77 (t, J = 4.7 Hz, 2 H), 6.82 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H), 6.95 (t, J = 8.7 Hz, 2 H), 7.02 (t, J = 8.6 Hz, 2 H), 7.15 (m, 2 H), 7.26 (m, 2 H), 7.89 (s, 1 H), 8.34 (dd, J_o = 1.6 Hz, J_m = 4.5 Hz, 2 H).

EXAMPLE 26

**4,6-Bis(4-fluorophenyl)-2-(4-methylsulfanylbenzyl)-5-(4-pyridyl)pyrazolo[3,4-
b]pyridine**

EXAMPLE 27

**4,6-Bis(4-fluorophenyl)-1-(4-methylsulfanylbenzyl)-5-(4-pyridyl)pyrazolo[3,4-
b]pyridine**

5

a) 1-Chloromethyl-4-(methylsulfanyl)benzene

A solution of thionyl chloride (0.2 mL, 3.2 mmol) in THF (9 mL) was added slowly to 4-(methylsulfanylphenyl)methanol (0.346g, 2.2 mmol) under argon atmosphere. The mixture was stirred at room temperature for 2 days. Brine (9 mL) was added and the phases were separated. The organic phase was dried over Na₂SO₄ and concentrated to dryness, to afford 0.37 g of the desired compound (yield: 95%).

10

b) Title compounds

In a volumetric flask, 4,6-bis(4-fluorophenyl)-5-(4-pyridyl)-1H-pyrazolo[3,4-b]pyridine (0.30 g, 78.0 mmol, obtained in example 1) and DMF (3.5 mL) were introduced under argon atmosphere. KOH (0.06 g, 1.1 mmol) was added followed by a solution of 1-chloromethyl-4-(methylsulfanyl)benzene (0.15 g, 0.9 mmol, obtained in section a) in DMF (0.4 mL). This was heated to 60 °C overnight. It was allowed to cool and concentrated. The residue was dissolved in a mixture of water and EtOAc. The two phases were separated. The organic phase was dried over Na₂SO₄ and concentrated to dryness. The crude product obtained was purified by chromatography on silica gel using hexane-EtOAc mixtures of increasing polarity as eluent, to afford 0.10 g of 4,6-bis(4-fluorophenyl)-2-(4-methylsulfanylbenzyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridine (yield: 25%) and 0.19 g of 4,6-bis(4-fluorophenyl)-1-(4-methylsulfanylbenzyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridine (yield: 47%).

20

25

Example 26: ¹H NMR (300 MHz, CDCl₃) δ (TMS): 2.50 (s, 3 H), 5.60 (s, 2 H), 6.84 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H), 6.92 (t, J = 8.8 Hz, 2 H), 7.00 (t, J = 8.7 Hz, 2 H), 7.05 (m, 2 H), 7.24 – 7.37 (complex signal, 6 H), 7.76 (s, 1 H), 8.34 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H).

30

Example 27: ¹H NMR (300 MHz, CDCl₃) δ (TMS): 2.49 (s, 3 H), 5.77 (s, 2 H), 6.85 (dd, J_o = 1.6 Hz, J_m = 4.4 Hz, 2 H), 6.97 (t, J = 8.8 Hz, 2 H), 7.03 (t, J = 8.7 Hz, 2 H), 7.15 (m, 2 H), 7.24 – 7.33 (complex signal, 4 H), 7.42 (d, J = 8.1 Hz, 2 H), 7.89 (s, 1 H), 8.36 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H).

EXAMPLE 28

2-[1-(*tert*-Butoxycarbonyl)piperidin-4-ylmethyl]-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine

EXAMPLE 29

5 1-[1-(*tert*-Butoxycarbonyl)piperidin-4-ylmethyl]-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine

a) (4-Piperidyl)methanol

To a suspension of LiAlH_4 (10.10 g, 0.266 mol) in THF (150 mL) at 0 °C, a solution of ethyl piperidine-4-carboxylate (18.13 g, 0.120 mol) in THF (300 mL) was added slowly. This was stirred at room temperature overnight. It was cooled with an ice bath and a mixture of water (14 mL) and THF (28 mL) was added slowly. Afterwards, a mixture of 15% aqueous NaOH (14 mL) and water (37 mL) was added followed by stirring at room temperature for 30 min. The precipitate obtained was filtered and the filtrate was concentrated, to afford 17.88 g of the desired compound (yield: quantitative).

b) [1-(*tert*-Butoxycarbonyl)piperidin-4-yl]methanol

Following a similar procedure to that described in Example 12 section a, but using (4-piperidyl)methanol (obtained in section a of this example), instead of 2-(4-piperidyl)ethanol, the title compound was obtained (yield: 77%).

20 c) [1-(*tert*-Butoxycarbonyl)piperidin-4-yl]methyl methanesulfonate

Following a similar procedure to that described in example 12 section b, but using [1-(*tert*-butoxycarbonyl)piperidin-4-yl]methanol (obtained in section b of this example) instead of 2-[1-(*tert*-butoxycarbonyl)piperidin-4-yl]ethanol, the title compound was obtained (yield: 71%).

25 d) Title compounds

Following a similar procedure to that described in examples 26 and 27, but using [1-(*tert*-butoxycarbonyl)piperidin-4-yl]methyl methanesulfonate (obtained in section c) instead of 1-chloromethyl-4-(methylsulfanyl)benzene, the title compounds were obtained.

30 Example 28: yield: 22%; ^1H NMR (300 MHz, CDCl_3) δ (TMS): 1.47 (s, 9 H), 1.63 (m, 2 H), 2.43 (m, 1 H), 2.72 (m, 2 H), 4.15 (m, 2 H), 4.33 (d, $J = 7.2$ Hz, 2 H), 6.86 (dd, $J_o = 1.5$ Hz, $J_m = 4.5$ Hz, 2 H), 6.93 (t, $J = 8.7$ Hz, 2 H), 7.04 (t, $J = 8.7$ Hz, 2

H), 7.17 (m, 2 H), 7.33 (m, 2 H), 7.77 (s, 1 H), 8.35 (dd, $J_o = 1.6$ Hz, $J_m = 4.4$ Hz, 2 H).

Example 29: yield: 71%; ^1H NMR (300 MHz, CDCl_3) δ (TMS): 1.20 – 1.8 (complex signal, 5 H), 1.49 (s, 9 H), 2.35 (m, 1 H), 2.74 (m, 2 H), 4.15 (m, 2 H), 4.54 (d, $J = 7.2$ Hz, 2 H), 6.86 (dd, $J_o = 1.5$ Hz, $J_m = 4.5$ Hz, 2 H), 6.97 (t, $J = 8.7$ Hz, 2 H), 7.04 (t, $J = 8.7$ Hz, 2 H), 7.18 (m, 2 H), 7.28 (m, 2 H), 7.90 (s, 1 H), 8.36 (dd, $J_o = 1.6$ Hz, $J_m = 4.5$ Hz, 2 H).

EXAMPLE 30

10 4,6-Bis(4-fluorophenyl)-2-[2-(morpholin-4-yl)ethyl]-5-(4-pyridyl)pyrazolo[3,4-
b]pyridine

EXAMPLE 31

4,6-Bis(4-fluorophenyl)-1-[2-(morpholin-4-yl)ethyl]-5-(4-pyridyl)pyrazolo[3,4-
b]pyridine

15 Following a similar procedure to that described in examples 26 and 27, but using
4-(2-chloroethyl)morpholine hydrochloride instead of 1-chloromethyl-4-
(methylsulfanyl)benzene and 2 equivalents of KOH, the title compounds were
obtained.

20 Example 30: yield: 10%; ^1H NMR (300 MHz, CDCl_3) δ (TMS): 2.55 (m, 4 H), 3.06
(t, $J = 6.4$ Hz, 2 H), 3.70 (m, 4 H), 4.58 (t, $J = 6.4$ Hz, 2 H), 6.86 (dd, $J_o = 1.6$ Hz,
 $J_m = 4.4$ Hz, 2 H), 6.93 (t, $J = 8.9$ Hz, 2 H), 7.04 (t, $J = 8.7$ Hz, 2 H), 7.16 (m, 2 H),
7.31 (m, 2 H), 7.88 (s, 1 H), 8.35 (dd, $J_o = 1.5$ Hz, $J_m = 4.5$ Hz, 2 H).

25 Example 31: yield: 24%; ^1H NMR (300 MHz, CDCl_3) δ (TMS): 2.63 (m, 4 H), 3.04
(t, $J = 6.9$ Hz, 2 H), 3.69 (m, 4 H), 4.77 (t, $J = 6.8$ Hz, 2 H), 6.86 (dd, $J_o = 1.5$ Hz,
 $J_m = 4.5$ Hz, 2 H), 6.96 (t, $J = 8.7$ Hz, 2 H), 7.05 (t, $J = 8.7$ Hz, 2 H), 7.18 (m, 2 H),
7.28 (m, 2 H), 7.90 (s, 1 H), 8.36 (dd, $J_o = 1.5$ Hz, $J_m = 4.5$ Hz, 2 H).

EXAMPLE 32

Ethyl 2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridin-2-
yl]acetate

EXAMPLE 33

30 Ethyl 2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridin-1-
yl]acetate

Following a similar procedure to that described in examples 26 and 27, but using ethyl bromoacetate instead of 1-chloromethyl-4-(methylsulfanyl)benzene, the title compounds were obtained.

Example 32: yield: 6%; ¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.32 (t, J = 7.2 Hz, 3 H), 4.32 (c, J = 7.2 Hz, 2 H), 5.27 (s, 2 H), 6.86 (dd, J_o = 1.6 Hz, J_m = 4.4 Hz, 2 H), 6.93 (t, J = 8.7 Hz, 2 H), 7.03 (t, J = 8.6 Hz, 2 H), 7.18 (m, 2 H), 7.32 (m, 2 H), 7.93 (s, 1 H), 8.35 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H).

Example 33: yield: 21%; ¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.32 (t, J = 7 Hz, 3 H), 4.31 (c, J = 6.9 Hz, 2 H), 5.42 (s, 2 H), 6.84 (dd, J_o = 1.6 Hz, J_m = 4.5 Hz, 2 H), 6.95 (t, J = 8.7 Hz, 2 H), 7.05 (t, J = 8.7 Hz, 2 H), 7.19 (m, 2 H), 7.28 (m, 2 H), 7.98 (s, 1 H), 8.36 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H).

EXAMPLE 34

Ethyl 3-[4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]propionate

EXAMPLE 35

Ethyl 3-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-1-yl]propionate

Following a similar procedure to that described in examples 26 and 27, but using ethyl 3-bromopropionate instead of 1-chloromethyl-4-(methylsulfanyl)benzene, the title compounds were obtained.

Example 34: yield: 5%; ¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.26 (t, J = 7.2 Hz, 3 H), 3.20 (t, J = 6.3 Hz, 2 H), 4.31 (c, J = 7.2 Hz, 2 H), 4.76 (t, J = 6.3 Hz, 2 H), 6.85 (dd, J_o = 1.6 Hz, J_m = 4.4 Hz, 2 H), 6.93 (t, J = 8.8 Hz, 2 H), 7.03 (t, J = 8.5 Hz, 2 H), 7.16 (m, 2 H), 7.32 (m, 2 H), 7.92 (s, 1 H), 8.34 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H).

Example 35: yield: 3%; ¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.25 (t, J = 7.2 Hz, 3 H), 3.10 (t, J = 7.2 Hz, 2 H), 4.17 (c, J = 7.1 Hz, 2 H), 4.94 (t, J = 7.0 Hz, 2 H), 6.85 (dd, J_o = 1.6 Hz, J_m = 4.6 Hz, 2 H), 6.96 (t, J = 8.6 Hz, 2 H), 7.04 (t, J = 8.6 Hz, 2 H), 7.18 (m, 2 H), 7.31 (m, 2 H), 7.89 (s, 1 H), 8.36 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H).

EXAMPLE 36

4,6-Bis(4-fluorophenyl)-2-(4-piperidyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine

To a solution of 2-[1-(*tert*-butoxycarbonyl)piperidin-4-yl]-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine (0.62 g, 1.1 mmol, obtained in example 20) in CH₂Cl₂ (19 mL) under argon atmosphere and cooled to 0 °C, trifluoroacetic acid (1.8 mL) was added. It was stirred at room temperature for 2.5 h. The solvent was evaporated. The residue was dissolved in CHCl₃ and washed with 1 N NaOH and brine. The organic phase was dried over Na₂SO₄ and concentrated to dryness, to afford 319 mg of the title compound (yield: 62%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.71 (broad s, NH + H₂O), 2.21 (m, 2 H), 2.34 (m, 2 H), 2.88 (m, 2 H), 3.35 (m, 2 H), 4.54 (m, 1 H), 6.87 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H), 6.95 (t, J = 8.7 Hz, 2 H), 7.06 (t, J = 8.6 Hz, 2 H), 7.18 (m, 2 H), 7.34 (m, 2 H), 7.88 (s, 1 H), 8.36 (dd, J_o = 1.6 Hz, J_m = 4.4 Hz, 2 H).

EXAMPLE 37

4,6-Bis(4-fluorophenyl)-1-(4-piperidyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine

Following a similar procedure to that described in example 36, but using 1-[1-(*tert*-butoxycarbonyl)piperidin-4-yl]-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine (obtained in example 21) instead of 2-[1-(*tert*-butoxycarbonyl)piperidin-4-yl]-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine, the title compound was obtained (yield: 29%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.62 (broad s, NH + H₂O), 2.13 (m, 2 H), 2.32 (m, 2 H), 2.92 (m, 2 H), 3.33 (m, 2 H), 4.52 (m, 1 H), 6.81 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H), 6.94 (t, J = 8.7 Hz, 2 H), 7.01 (t, J = 8.6 Hz, 2 H), 7.14 (m, 2 H), 7.27 (m, 2 H), 7.86 (s, 1 H), 8.32 (dd, J_o = 1.4 Hz, J_m = 4.4 Hz, 2 H).

EXAMPLE 38

4,6-Bis(4-fluorophenyl)-2-(4-piperidylmethyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine

Following a similar procedure to that described in example 36, but using 2-[1-(*tert*-butoxycarbonyl)piperidin-4-ylmethyl]-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine (obtained in example 28) instead of 2-[1-(*tert*-butoxycarbonyl)piperidin-4-yl]-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine, the title compound was obtained (yield: 13%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.30 - 1.80 (complex signal, 4 H), 1.63 (broad s, NH + H₂O), 2.38 (m, 1 H), 2.64 (m, 2 H), 3.14 (m, 2 H), 4.32 (d, J = 7.2 Hz), 6.86 (dd, J_o = 1.6 Hz, J_m = 4.4 Hz, 2 H), 6.93 (t, J = 8.8 Hz, 2 H), 7.04 (t, J = 8.6 Hz, 2 H).

H), 7.18 (m, 2 H), 7.33 (m, 2 H), 7.78 (s, 1 H), 8.35 (dd, $J_o = 1.8$ Hz, $J_m = 4.5$ Hz, 2 H).

EXAMPLE 39

4,6-Bis(4-fluorophenyl)-1-(4-piperidylmethyl)-5-(4-pyridyl)pyrazolo[3,4- b]pyridine

5

Following a similar procedure to that described in example 36, but using 1-[1-(*tert*-butoxycarbonyl)piperidin-4-ylmethyl]-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-
b]pyridine (obtained in example 29) instead of 2-[1-(*tert*-butoxycarbonyl)piperidin-4-yl]-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-
b]pyridine, the title compound was obtained (yield: 12%).

10

^1H NMR (300 MHz, CDCl_3) δ (TMS): 1.80 – 2.10 (complex signal, 4 H), 2.47 (m, 1 H), 3.0 (m, 3 H), 3.50 (m, 2 H), 4.63 (d, $J = 6.6$, 2 H), 7.04 (t, $J = 8.4$ Hz, 2 H), 7.12 – 7.16 (complex signal, 4 H), 7.22 (m, 2 H), 7.31 (m, 2 H), 7.95 (s, 1 H), 8.58 (d, $J = 6.6$, 2 H).

15

EXAMPLE 40

4,6-Bis(6-chloropyridin-3-yl)-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine

Following a similar procedure to that described in example 1 method A, but using 1-(6-chloropyridin-3-yl)-2-(4-pyridyl)ethanone (obtained in reference example 9) instead of 1-(4-fluorophenyl)-2-(4-pyridyl)ethanone, the title compound was
obtained (yield: 17%).

20

^1H NMR (300 MHz, CDCl_3) δ (TMS): 6.90 (dd, $J_o = 1.5$ Hz, $J_m = 4.5$ Hz, 2 H), 7.21 (d, $J = 8.4$ Hz, 1 H), 7.30 (d, $J = 8.4$ Hz, 1 H), 7.38 (dd, $J_o = 2.5$ Hz, $J_m = 8.3$ Hz, 1 H), 7.49 (dd, $J_o = 2.4$ Hz, $J_m = 8.4$ Hz, 1 H), 8.00 (s, 1 H), 8.36 (d, $J = 2.4$ Hz, 1 H), 8.46 – 8.48 (complex signal, 3 H), 11.34 (broad s, 1 H).

25

EXAMPLE 41

4,6-Bis(4-fluorophenyl)-3-methyl-2-(4-piperidyl)-5-(4-pyridyl)pyrazolo[3,4- b]pyridine

Following a similar procedure to that described in example 36, but using 2-[1-(*tert*-butoxycarbonyl)piperidin-4-yl]-4,6-bis(4-fluorophenyl)-3-methyl-5-(4-pyridyl)pyrazolo[3,4-
b]pyridine (obtained in example 14) instead of 2-[1-(*tert*-butoxycarbonyl)piperidin-4-yl]-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-
b]pyridine, the title compound was obtained (yield: 73%).

30

¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.55 (broad s, NH + H₂O), (s, 3 H), 2.35 (m, 2 H), 3.34 (m, 2 H), 3.85 (m, 2 H), 4.73 (m, 2 H), 4.75 (m, 1 H), 6.78 (d, J = 6.3 Hz, 2 H), 6.89 (t, J = 8.6 Hz, 2 H), 7.03 – 7.19 (complex signal, 4 H), 7.29 (m, 2 H), 8.27 (d, J = 6.0 Hz, 2 H).

5

EXAMPLE 42**4,6-Bis(4-fluorophenyl)-3-methyl-1-(4-piperidyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridine**

Following a similar procedure to that described in example 36, but using 1-[1-(*tert*-butoxycarbonyl)piperidin-4-yl]-4,6-bis(4-fluorophenyl)-3-methyl-5-(4-pyridyl)pyrazolo[3,4-b]pyridine (obtained in example 15) instead of 2-[1-(*tert*-butoxycarbonyl)piperidin-4-yl]-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridine, the title compound was obtained (yield: 59%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.55 (broad s, NH + H₂O), 2.01 (s, 3 H), 2.07 (m, 2 H), 2.29 (m, 2 H), 2.90 (m, 2 H), 3.32 (m, 2 H), 4.75 (m, 1 H), 6.78 (dd, J_o = 1.6 Hz, J_m = 4.4 Hz, 2 H), 6.93 (t, J = 8.7 Hz, 2 H), 6.99 (t, J = 8.7 Hz, 2 H), 7.08 (m, 2 H), 7.30 (m, 2 H), 8.27 (dd, J_o = 1.6 Hz, J_m = 4.4 Hz, 2 H).

15

EXAMPLE 43**4,6-Bis(4-fluorophenyl)-2-[2-(4-piperidyl)ethyl]-5-(4-pyridyl)pyrazolo[3,4-b]pyridine**

Following a similar procedure to that described in example 36, but using 2-[2-[1-(*tert*-butoxycarbonyl)piperidin-4-yl]ethyl]-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridine (obtained in example 12) instead of 2-[1-(*tert*-butoxycarbonyl)piperidin-4-yl]-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridine, the title compound was obtained (yield: 88%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.00 – 1.60 (complex signal, 3 H), 1.66 (s, NH + H₂O), 1.74 (m, 2 H), 2.03 (m, 2 H), 2.60 (m, 2 H), 3.10 (m, 2 H), 4.85 (t, J = 7.2 Hz, 2 H), 6.82 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H), 6.89 (t, J = 8.7 Hz, 2 H), 7.00 (t, J = 8.7 Hz, 2 H), 7.13 (m, 2 H), 7.30 (m, 2 H), 7.76 (s, 1 H), 8.31 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H).

25

30

EXAMPLE 44**2-[4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridin-2-yl]acetic acid**

To a solution of ethyl 2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridin-2-yl]acetate (0.09 g, 0.2 mmol, obtained in example 32) in EtOH (4.2 mL), a

solution of KOH (0.09 g, 2.5 mmol) in water (0.5 mL) was added. This was heated to reflux for 1 h. It was allowed to cool and concentrated. The residue was dissolved in a mixture of EtOAc and water. The phases were separated. The aqueous phase was acidified and extracted with EtOAc. The combined organic phases were dried over Na₂SO₄ and concentrated to dryness, to afford 57 mg of title compound (yield: 66%).

¹H NMR (300 MHz, CDCl₃ + CD₃OD) δ (TMS): 4.00 (broad s, 1H + CD₃OD), 5.28 (broad s, 2 H), 6.92 – 7.05 (complex signal, 6 H), 7.15 – 7.30 (complex signal, 4 H), 8.02 (broad s, 1 H), 8.26 (m, 2 H).

10

EXAMPLE 45

2-[4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-1-yl]acetic acid

Following a similar procedure to that described in example 44, but using ethyl 2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-1-yl]acetate (obtained in example 33) instead of ethyl 2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]acetate, the title compound was obtained (yield: 96%).

15

¹H NMR (300 MHz, CDCl₃ + CD₃OD) δ (TMS): 3.88 (broad s, 1H + CD₃OD), 5.32 (s, 2 H), 6.83 (dd, *J*_o = 1.5 Hz, *J*_m = 4.5 Hz, 2 H), 6.88 (t, *J* = 8.7 Hz, 2 H), 6.98 (t, *J* = 8.6 Hz, 2 H), 7.11 (m, 2 H), 7.20 (m, 2 H), 7.90 (s, 1 H), 8.21 (dd, *J*_o = 1.5 Hz, *J*_m = 4.5 Hz, 2 H).

20

EXAMPLE 46

3-[4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]propionic acid

Following a similar procedure to that described in example 44, but using ethyl 3-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]propionate (obtained in example 34) instead of ethyl 2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]acetate, the title compound was obtained (yield: 69%).

25

¹H NMR (300 MHz, CDCl₃ + CD₃OD) δ (TMS): 3.08 (t, *J* = 6.3 Hz, 2 H), 4.50 (broad s, 1H + CD₃OD), 4.72 (t, *J* = 6.3 Hz, 2 H), 6.88 (dd, *J*_o = 1.6 Hz, *J*_m = 4.6 Hz, 2 H), 6.90 (t, *J* = 8.7 Hz, 2 H), 6.99 (t, *J* = 8.7 Hz, 2 H), 7.15 (m, 2 H), 7.23 (m, 2 H), 8.03 (s, 1 H), 8.21 (dd, *J*_o = 1.5 Hz, *J*_m = 4.5 Hz, 2 H).

30

EXAMPLE 47

3-[4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-1-yl]propionic acid

Following a similar procedure to that described in example 44, but using ethyl 3-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-1-yl]propionate (obtained in example 35) instead of ethyl 2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]acetate, the title compound was obtained (yield: 88%).

¹H NMR (300 MHz, CDCl₃ + CD₃OD) δ (TMS): 3.02 (t, J = 7.2 Hz, 2 H), 4.50 (broad s, 1H + CD₃OD), 4.86 (t, J = 7.3 Hz, 2 H), 6.87 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H), 6.91 (t, J = 8.7 Hz, 2 H), 7.00 (t, J = 8.7 Hz, 2 H), 7.12 (m, 2 H), 7.24 (m, 2 H), 7.84 (s, 1 H), 8.21 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H).

EXAMPLE 48

2-[4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]-1-(morpholin-4-yl)ethanone

To a solution of 2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]acetic acid (0.05 g, 0.1 mmol, obtained in example 44) in DMF (1 mL), *N,N'*-dicyclohexylcarbodiimide (0.02 g, 0.1 mmol) and 1-hydroxybenzotriazole (0.02 g, 0.1 mmol) were added under argon atmosphere. The mixture was stirred for 15 min and morpholine (0.01 g, 0.1 mmol) was added. This was stirred at room temperature for 2 days. EtOAc was added and the mixture was filtered. The filtrate was washed with saturated NaHCO₃. The organic phase was dried over Na₂SO₄ and concentrated to dryness. The crude product obtained was purified by chromatography on silica gel using EtOAc-MeOH mixtures of increasing polarity as eluent, to afford 18 mg of title compound (yield: 35%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 3.66 – 3.76 (complex signal, 8 H), 5.30 (s, 2 H), 6.82 (d, J = 5.7 Hz, 2 H), 6.91 (t, J = 8.8 Hz, 2 H), 6.99 (t, J = 8.7 Hz, 2 H), 7.14 (m, 2 H), 7.28 (m, 2 H), 7.99 (s, 1 H), 8.32 (d, J = 6.0 Hz, 2 H).

EXAMPLE 49

2-[4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-1-yl]acetamide

Following a similar procedure to that described in example 48, but using 2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-1-yl]acetic acid (obtained in example 45) instead of 2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-

b]pyridin-2-yl]acetic acid, and aqueous NH₃ instead of morpholine, the title compound was obtained (yield: 36%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 5.35 (s, 2 H), 5.56 (broad s, 1 H), 6.15 (broad s, 1 H), 6.85 (d, J = 5.4 Hz, 2 H), 6.98 (t, J = 8.8 Hz, 2 H), 7.08 (t, J = 8.6 Hz, 2 H),
5 7.17 (m, 2 H), 7.29 (m, 2 H), 8.01 (s, 1 H), 8.38 (d, J = 5.4 Hz, 2 H).

EXAMPLE 50

2-[4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridin-1-yl]-1-(morpholin-4-yl)ethanone

Following a similar procedure to that described in example 48, but using 2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridin-1-yl]acetic acid (obtained in
10 example 45) instead of 2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridin-2-yl]acetic acid, the title compound was obtained (yield: 60%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 3.67 – 3.70 (complex signal, 4 H), 3.75 – 3.81 (complex signal, 4 H), 5.50 (s, 2 H), 6.83 (dd, J_o = 1.6 Hz, J_m = 4.4 Hz, 2 H), 6.95
15 (t, J = 8.7 Hz, 2 H), 7.04 (t, J = 8.7 Hz, 2 H), 7.17 (m, 2 H), 7.27 (m, 2 H), 7.99 (s, 1 H), 8.35 (dd, J_o = 1.6 Hz, J_m = 4.4 Hz, 2 H)

EXAMPLE 51

3-[4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridin-2-yl]-1-(morpholin-4-yl)propan-1-one

20 Following a similar procedure to that described in example 48, but using 3-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridin-2-yl]propionic acid (obtained in example 46) instead of 2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridin-2-yl]acetic acid, the title compound was obtained (yield: 64%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 3.20 (t, J = 6.0 Hz, 2 H), 3.45 (m, 2 H), 3.55 –
25 3.63 (complex signal, 6 H), 4.79 (t, J = 6.1 Hz, 2 H), 6.81 (dd, J_o = 1.6 Hz, J_m = 4.4 Hz, 2 H), 6.90 (t, J = 8.7 Hz, 2 H), 6.99 (t, J = 8.7 Hz, 2 H), 7.13 (m, 2 H), 7.30 (m, 2 H), 7.94 (s, 1 H), 8.31 (dd, J_o = 1.6 Hz, J_m = 4.4 Hz, 2 H)

EXAMPLE 52

3-[4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridin-1-yl]-N-propylpropionamide

30

Following a similar procedure to that described in example 48, but using 3-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridin-1-yl]propionic acid (obtained in example 47) instead of 2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-

b]pyridin-2-yl]acetic acid, and propylamine instead of morpholine, the title compound was obtained (yield: 76%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 0.84 (t, J = 7.3 Hz, 3 H), 1.42 (m, 2 H), 2.98 (t, J = 6.7 Hz, 2 H), 3.18 (m, 2 H), 4.95 (t, J = 6.7 Hz, 2 H), 6.05 (m, NH), 6.84 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H), 6.96 (t, J = 8.7 Hz, 2 H), 7.04 (t, J = 8.7 Hz, 2 H), 7.16 (m, 2 H), 7.32 (m, 2 H), 7.89 (s, 1 H), 8.36 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H).

EXAMPLE 53

3-[4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-1-yl]-1-(morpholin-4-yl)propan-1-one

10 Following a similar procedure to that described in example 48, but using 3-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-1-yl]propionic acid (obtained in example 47) instead of 2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]acetic acid, the title compound was obtained (yield: 72%).

15 ¹H NMR (300 MHz, CDCl₃) δ (TMS): 3.13 (t, J = 7.3 Hz, 2 H), 3.49 (m, 2 H), 3.63 – 3.67 (complex signal, 6 H), 4.99 (t, J = 7.3 Hz, 2 H), 6.84 (dd, J_o = 1.4 Hz, J_m = 4.4 Hz, 2 H), 6.96 (t, J = 8.7 Hz, 2 H), 7.04 (t, J = 8.7 Hz, 2 H), 7.16 (m, 2 H), 7.30 (m, 2 H), 7.89 (s, 1 H), 8.36 (dd, J_o = 1.6 Hz, J_m = 4.4 Hz, 2 H)

EXAMPLE 54

4,6-Bis(4-fluorophenyl)-2-(4-methylsulfanylphenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine

EXAMPLE 55

4,6-Bis(4-fluorophenyl)-1-(4-methylsulfanylphenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine

25 In a volumetric flask molecular sieves of 4 Å (1 g, previously dried for 3 h at 200 °C under vacuum), 4,6-bis(4-fluorophenyl)-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine (0.30 g, 0.8 mmol, obtained in example 1), (4-methylsulfanylphenyl)boronic acid (0.26 g, 1.6 mmol), copper II acetate (0.28 g, 1.6 mmol), pyridine (0.12 g, 1.6 mmol), triethylamine (0.16 g, 1.6 mmol) and CH₂Cl₂ (22 mL) were introduced under argon atmosphere. This was stirred at room temperature for 2 days. It was
30 filtered through celite and concentrated. The crude product obtained was purified by chromatography on silica gel using hexane-EtOAc mixtures of increasing polarity as eluent, to afford 40 mg of 4,6-bis(4-fluorophenyl)-2-(4-methylsulfanylphenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine (yield: 10%) and 90 mg

of 4,6-bis(4-fluorophenyl)-1-(4-methylsulfonylphenyl)-5-(4-pyridyl) pyrazolo[3,4-*b*]pyridine (yield: 23%).

Example 54: ¹H NMR (300 MHz, CDCl₃) δ (TMS): 2.46 (s, 3 H), 6.76 (dd, *J*_o = 1.6 Hz, *J*_m = 4.4 Hz, 2 H), 6.83 (t, *J* = 8.7 Hz, 2 H), 6.95 (t, *J* = 8.6 Hz, 2 H), 7.11 (m, 2 H), 7.26 (m, 2 H), 7.29 (d, *J* = 8.7 Hz, 2 H), 7.83 (d, *J* = 8.7 Hz, 2 H), 8.16 (s, 1 H), 8.26 (dd, *J*_o = 1.6 Hz, *J*_m = 4.4 Hz, 2 H).

Example 55: ¹H NMR (300 MHz, CDCl₃) δ (TMS): 2.46 (s, 3 H), 6.76 (dd, *J*_o = 1.5 Hz, *J*_m = 4.5 Hz, 2 H), 6.86 (t, *J* = 8.7 Hz, 2 H), 6.96 (t, *J* = 8.7 Hz, 2 H), 7.08 (m, 2 H), 7.22 (m, 2 H), 7.34 (d, *J* = 9.0 Hz, 2 H), 7.94 (s, 1 H), 8.23 (d, *J* = 8.7 Hz, 2 H), 8.26 (dd, *J*_o = 1.5 Hz, *J*_m = 4.5 Hz, 2 H).

EXAMPLE 56

4,6-Bis(4-fluorophenyl)-2-(4-methylsulfinylphenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine

EXAMPLE 57

4,6-Bis(4-fluorophenyl)-2-(4-methylsulfonylphenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine

To a solution of 4,6-bis(4-fluorophenyl)-2-(4-methylsulfonylphenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine (0.09 g, 0.2 mmol, obtained in example 54) in CH₂Cl₂ (3.5 mL), 3-chloroperbenzoic acid (0.04 g, 0.2 mmol) was added under argon atmosphere and stirred for 2 h at room temperature. CHCl₃ was added and washed with saturated NaHCO₃. The organic phase was dried over Na₂SO₄ and concentrated to dryness. The crude product obtained was purified by chromatography on silica gel using increasing polarity hexane-EtOAc mixtures as eluent, to afford 15 mg of 4,6-bis(4-fluorophenyl)-2-(4-methylsulfinylphenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine (yield: 16%) and 10 mg of 4,6-bis(4-fluorophenyl)-2-(4-methylsulfonylphenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine (yield: 11%).

Example 56: ¹H NMR (300 MHz, CDCl₃) δ (TMS): 2.83 (s, 3 H), 6.90 (dd, *J*_o = 1.5 Hz, *J*_m = 4.5 Hz, 2 H), 6.96 (t, *J* = 8.7 Hz, 2 H), 7.08 (t, *J* = 8.7 Hz, 2 H), 7.24 (m, 2 H), 7.36 (m, 2 H), 7.75 (H_A from an AB system, *J* = 8.9 Hz, 2 H), 8.32 (H_B from an AB system, *J* = 8.9 Hz, 2 H), 8.38 (s, 1 H), 8.39 (dd, *J*_o = 1.5 Hz, *J*_m = 4.5 Hz, 2 H).

Example 57: ¹H NMR (300 MHz, CDCl₃) δ (TMS): 3.16 (s, 3 H), 6.89 (dd, *J*_o = 1.6 Hz, *J*_m = 4.4 Hz, 2 H), 6.97 (t, *J* = 8.7 Hz, 2 H), 7.09 (t, *J* = 8.7 Hz, 2 H), 7.24 (m, 2

H), 7.36 (m, 2 H), 8.05 (H_A from an AB system, $J = 9.0$ Hz, 2 H), 8.38 (H_B from an AB system, $J = 9.0$ Hz, 2 H), 8.39 (dd, $J_o = 1.6$ Hz, $J_m = 4.6$ Hz, 2 H), 8.41 (s, 1 H).

EXAMPLE 58

4,6-Bis(4-fluorophenyl)-1-(4-methylsulfinylphenyl)-5-(4-pyridyl)pyrazolo[3,4- b]pyridine

Following a similar procedure to that described in example 56, but using 4,6-bis(4-fluorophenyl)-1-(4-methylsulfinylphenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine (obtained in example 55) instead of 4,6-bis(4-fluorophenyl)-2-(4-methylsulfinylphenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine, the title compound was obtained (yield: 70%).

1H NMR (300 MHz, $CDCl_3$) δ (TMS): 2.82 (s, 3 H), 6.90 (dd, $J_o = 1.8$ Hz, $J_m = 4.5$ Hz, 2 H), 7.00 (t, $J = 8.7$ Hz, 2 H), 7.09 (t, $J = 8.6$ Hz, 2 H), 7.20 (m, 2 H), 7.35 (m, 2 H), 7.86 (dd, $J_o = 2.1$ Hz, $J_m = 6.9$ Hz, 2 H), 8.10 (s, 1 H), 8.41 (dd, $J_o = 1.5$ Hz, $J_m = 4.5$ Hz, 2 H), 8.69 (dd, $J_o = 1.8$ Hz, $J_m = 6.9$ Hz, 2 H).

EXAMPLE 59

4,6-Bis(4-fluorophenyl)-2-(4-methylsulfinylbenzyl)-5-(4-pyridyl)pyrazolo[3,4- b]pyridine

EXAMPLE 60

4,6-Bis(4-fluorophenyl)-2-(4-methylsulfonylbenzyl)-5-(4-pyridyl)pyrazolo[3,4- b]pyridine

Following a similar procedure to that described in example 56, but using 4,6-bis(4-fluorophenyl)-2-(4-methylsulfonylbenzyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine (obtained in example 26) instead of 4,6-bis(4-fluorophenyl)-2-(4-methylsulfinylphenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine, the title compounds were obtained.

Example 59: yield: 48%; 1H NMR (300 MHz, $CDCl_3$) δ (TMS): 2.74 (s, 3 H), 5.70 (s, 2 H), 6.85 (dd, $J_o = 1.5$ Hz, $J_m = 4.5$ Hz, 2 H), 6.93 (t, $J = 8.8$ Hz, 2 H), 7.02 (t, $J = 8.7$ Hz, 2 H), 7.16 (m, 2 H), 7.32 (m, 2 H), 7.59 (H_A from an AB system, $J = 8.2$ Hz, 2 H), 7.68 (H_B from an AB system, $J = 8.2$ Hz, 2 H), 7.87 (s, 1 H), 8.39 (dd, $J_o = 1.5$ Hz, $J_m = 4.5$ Hz, 2 H).

Example 60: yield: 16%; 1H NMR (300 MHz, $CDCl_3$) δ (TMS): 3.07 (s, 3 H), 5.73 (s, 2 H), 6.85 (dd, $J_o = 1.6$ Hz, $J_m = 4.4$ Hz, 2 H), 6.93 (t, $J = 8.7$ Hz, 2 H), 7.03 (t, $J = 8.8$ Hz, 2 H), 7.16 (m, 2 H), 7.32 (m, 2 H), 7.61 (H_A from an AB system, $J = 8.7$

Hz, 2 H), 7.89 (s, 1 H), 7.97 (H_B from an AB system, $J = 8.7$ Hz, 2 H), 8.36 (dd, $J_o = 1.8$ Hz, $J_m = 4.5$ Hz, 2 H).

EXAMPLE 61

4,6-Bis(4-fluorophenyl)-1-(4-methylsulfinylbenzyl)-5-(4-pyridyl)pyrazolo[3,4-
b]pyridine

EXAMPLE 62

4,6-Bis(4-fluorophenyl)-1-(4-methylsulfonylbenzyl)-5-(4-pyridyl)pyrazolo[3,4-
b]pyridine

Following a similar procedure to that described in example 56, but using 4,6-bis(4-fluorophenyl)-1-(4-methylsulfonylbenzyl)-5-(4-pyridyl)pyrazolo[3,4-
b]pyridine (obtained in example 27) instead of 4,6-bis(4-fluorophenyl)-2-(4-methylsulfonylphenyl)-5-(4-pyridyl)pyrazolo[3,4-
b]pyridine, the title compounds were obtained.

Example 61: yield: 69%; 1H NMR (300 MHz, $CDCl_3$) δ (TMS): 2.71 (s, 3 H), 5.84 (s, 2 H), 6.83 (dd, $J_o = 1.5$ Hz, $J_m = 4.5$ Hz, 2 H), 6.94 (t, $J = 8.7$ Hz, 2 H), 7.01 (t, $J = 8.6$ Hz, 2 H), 7.13 (m, 2 H), 7.25 (m, 2 H), 7.61 (H_A from an AB system, $J = 8.4$ Hz, 2 H), 7.92 (H_B from an AB system, $J = 8.4$ Hz, 2 H), 7.91 (s, 1 H), 8.34 (dd, $J_o = 1.5$ Hz, $J_m = 4.5$ Hz, 2 H).

Example 62: yield: 4%; 1H NMR (300 MHz, $CDCl_3$) δ (TMS): 3.03 (s, 3 H), 5.87 (s, 2 H), 6.83 (d, $J = 6.0$ Hz, 2 H), 6.94 (t, $J = 8.8$ Hz, 2 H), 7.02 (t, $J = 8.7$ Hz, 2 H), 7.14 (m, 2 H), 7.26 (m, 2 H), 7.60 (H_A from an AB system, $J = 8.2$ Hz, 2 H), 7.91 (s, 1 H), 7.93 (H_B from an AB system, $J = 8.2$ Hz, 2 H), 8.34 (d, $J = 6.3$ Hz, 2 H).

EXAMPLE 63

3-Chloro-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)-1*H*-pyrazolo[3,4-
b]pyridine

To a solution of 4,6-bis(4-fluorophenyl)-5-(4-pyridyl)-1*H*-pyrazolo[3,4-
b]pyridine (0.20 g, 0.5 mmol, obtained in example 1) in DMF (5 mL), *N*-chlorosuccinimide (0.10 g, 0.8 mmol) was added under argon atmosphere and the mixture was heated to 60 °C for 5 h. It was washed with 1 N NaOH and extracted with $CHCl_3$ and EtOAc. The combined organic phases were dried over Na_2SO_4 and concentrated to dryness. The crude product obtained was purified by chromatography on silica gel using hexane-EtOAc mixtures of increasing polarity as eluent, to afford 171 mg of title compound (yield: 79%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.61 (broad s, NH + H₂O), 6.82 (dd, J_o = 1.6 Hz, J_m = 4.4 Hz, 2 H), 6.98 (t, J = 8.7 Hz, 2 H), 7.03 (t, J = 8.7 Hz, 2 H), 7.13 (m, 2 H), 7.29 (m, 2 H), 8.35 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H).

EXAMPLE 64

5 3-Bromo-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)-1H-pyrazolo[3,4-b]pyridine

To a suspension of 4,6-bis(4-fluorophenyl)-5-(4-pyridyl)-1H-pyrazolo[3,4-b]pyridine (1.00 g, 2.6 mmol, obtained in example 1) in CHCl₃ (10 mL), a solution of Br₂ (0.69 g, 4.3 mmol) in CHCl₃ (3 mL) was added under argon atmosphere. Acetonitrile (4 mL) was added and the mixture was stirred at room temperature for 2 days. The residue was concentrated, dissolved in CHCl₃ and washed with 1 N NaOH. A precipitate was formed, which was filtered and dissolved with MeOH. The solution was concentrated and washed with 1 N NaOH. It was extracted with EtOAc, dried over Na₂SO₄ and concentrated. This was treated with diethyl ether, the solvent was decanted and the product obtained was dried to afford 1.16 g of the title compound (yield: 97%).

¹H NMR (300 MHz, CD₃OD) δ (TMS): 4.78 (broad s, NH + CD₃OD), 6.89 (t, J = 8.8 Hz, 2 H), 6.93 – 6.99 (complex signal, 4 H), 7.14 (m, 2 H), 7.27 (m, 2 H), 8.12 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H).

EXAMPLE 65

20 4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)-1H-pyrazolo[3,4-b]pyridine-3-carbonitrile

In a volumetric flask, 3-bromo-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)-1H-pyrazolo[3,4-b]pyridine (0.20 g, 0.4 mmol, obtained in example 64), copper cyanide (I) (0.05 g, 0.6 mmol) and anhydrous 1-methyl-2-pyrrolidone (1 mL) were introduced under argon atmosphere and heated to reflux for 2 h. This was poured into a 10% aqueous solution of ethylenediamine (4 mL) and extracted with CHCl₃. Brine was added to the aqueous phase and it was extracted with EtOAc. The combined organic phases were dried over Na₂SO₄ and concentrated to dryness. The crude product obtained was purified by chromatography on silica gel using hexane-EtOAc mixtures of increasing polarity as eluent, to afford the title compound in a solid form (yield: quantitative).

¹H NMR (300 MHz, CDCl₃+ CD₃OD) δ (TMS): 4.00 (broad s, NH + CD₃OD), 6.80 – 7.40 (complex signal, 12 H), 8.25 (m, 2 H).

EXAMPLE 66**3-Bromo-4,6-bis(4-fluorophenyl)-1-methyl-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine**

Following a similar procedure to that described in example 64, but using 4,6-bis(4-fluorophenyl)-1-methyl-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine (obtained in example 9) instead of 4,6-bis(4-fluorophenyl)-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine, the title compound was obtained (yield: 15%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 4.21 (s, 3 H), 6.78 (d, J = 6.0 Hz, 2 H), 6.93 – 7.02 (complex signal, 4 H), 7.08 (m, 2 H), 7.27 (m, 2 H), 8.29 (d, J = 6.0 Hz, 2 H).

EXAMPLE 67**4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine-3-carboxamide**

A solution of KOH (0.07 g, 1.3 mmol) in ^tBuOH (2.5 mL) was added under argon atmosphere to 4,6-bis(4-fluorophenyl)-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine-3-carbonitrile (0.05 g, 0.1 mmol, obtained in example 65) and heated to reflux overnight. Water and EtOAc were added and the phases were separated. The aqueous phase was extracted with EtOAc. The combined organic phases were dried over Na₂SO₄ and concentrated to dryness. The crude product obtained was purified by chromatography on silica gel using hexane-EtOAc mixtures of increasing polarity as eluent, to afford 15 mg of the title compound in solid form (yield: 28%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.56 (broad s, NH₂ + H₂O), 6.80 (d, J = 4.5 Hz, 2 H), 6.90 – 7.10 (complex signal, 6 H), 7.30 (m, 2 H), 8.30 (d, J = 4.5 Hz, 2 H).

EXAMPLE 68**3-Aminomethyl-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine**

LiAlH₄ (0.06 g, 1.5 mmol) and anhydrous diethyl ether (2 mL) were introduced into a volumetric flask. The mixture was cooled with an ice bath and a solution of 4,6-bis(4-fluorophenyl)-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine-3-carbonitrile (0.15 g, 0.4 mmol, obtained in example 65) in diethyl ether (1 mL) was added dropwise. THF (2 mL) was added and the mixture was stirred at room temperature overnight. It was cooled with an ice bath and successively water (0.1 mL), THF (0.2 mL), a

15% aqueous NaOH (0.1 mL) and water (0.3 mL) were added. The precipitate obtained was filtered and washed with THF. The solvent of the filtrate was evaporated. The residue obtained was purified by chromatography on silica gel using hexane-EtOAc mixtures of increasing polarity as eluent, to afford 71 mg of the title compound in solid form (yield: 47%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 2.10 (broad s, NH₂ + H₂O) 3.62 (s, 2 H), 6.80 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H), 6.93 (t, J = 8.7 Hz, 2 H), 7.03 (t, J = 8.6 Hz, 2 H), 7.11 (m, 2 H), 7.25 (m, 2 H), 8.30 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H).

EXAMPLE 69

10 **4,6-Bis(4-fluoro-3-nitrophenyl)-5-(4-pyridyl)-1H-pyrazolo[3,4-b]pyridine**

To a solution of 4,6-bis(4-fluorophenyl)-5-(4-pyridyl)-1H-pyrazolo[3,4-b]pyridine (0.20 g, 0.5 mmol, obtained in example 1) in conc. H₂SO₄ (3 mL), 65% HNO₃ (0.1 mL, 0.2 mmol) was added under argon atmosphere. This was heated to 90 °C for 30 min. It was cooled with an ice bath and adjusted to pH = 8 with 1 N NaOH. It was extracted with EtOAc. The organic phase was dried over Na₂SO₄ and concentrated to dryness. The crude product was purified by chromatography on silica gel using hexane-EtOAc mixtures of increasing polarity as eluent, to afford 69 mg of the title compound in solid form (yield: 28%).

¹H NMR (300 MHz, CDCl₃ + CD₃OD) δ (TMS): 4.28 (s, NH + CD₃OD), 7.03 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H), 7.22 (dd, J_o = 2.7 Hz, J_m = 10.5 Hz, 1 H), 7.36 (dd, J_o = 2.6 Hz, J_m = 10.4 Hz, 1 H), 7.53 (m, 2 H), 7.99 - 8.01 (complex signal, 2 H), 8.13 (m, 1 H), 8.41 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H).

EXAMPLE 70

3-Amino-6-(4-fluorophenyl)-5-(4-pyridyl)-1H-pyrazolo[3,4-b]pyridine

25 Following a similar procedure to that described in reference example 15 section c, but using 6-chloro-2-(4-fluorophenyl)-3,4'-bipyridine-5-carbonitrile (obtained in reference example 12) instead of 3-(1-benzylpiperidin-4-yl)-3-oxopropiononitrile, the title compound was obtained (yield: 41%).

¹H NMR (300 MHz, CD₃OD) δ (TMS): 4.38 (s, NH₂ + CD₃OD), 7.05 (t, J = 8.8 Hz, 2 H), 7.26 (dd, J_o = 1.6 Hz, J_m = 4.6 Hz, 2 H), 7.39 (m, 2 H), 8.27 (s, 1 H), 8.42 (dd, J_o = 1.4 Hz, J_m = 4.6 Hz, 2 H).

EXAMPLE 71

3-Amino-6-(4-fluorophenyl)-1-methyl-5-(4-pyridyl)pyrazolo[3,4-b]pyridine

Following a similar procedure to that described in reference example 15 section c, but using 6-chloro-2-(4-fluorophenyl)-3,4'-bipyridine-5-carbonitrile (obtained in reference example 12) instead of 3-(1-benzylpiperidin-4-yl)-3-oxopropiononitrile and methylhydrazine instead of hydrazine monohydrate, the title compound was
5 obtained (yield: 70%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 4.00 (s, 3 H), 4.17 (broad s, 2 H), 6.98 (t, J = 8.7 Hz, 2 H), 7.09 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H), 7.37 (m, 2 H), 7.90 (s, 1 H), 8.50 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H).

EXAMPLE 72

10 4-[6-(4-Fluorophenyl)-5-(4-pyridyl)-1H-pyrazolo[3,4-b]pyridin-4-yl]phenol

To a solution of 1-(4-fluorophenyl)-2-(4-pyridyl)ethanone (0.30 g, 1.4 mmol, obtained in reference example 1) in 2-methoxyethanol (2 mL), 3-amino-2H-pyrazole (0.13 g, 1.5 mmol), 4-hydroxybenzaldehyde (0.17 g, 1.4 mmol), 2-methoxyethanol (2 mL) and 37% HCl (0.04 g, 0.4 mmol) were added under argon
15 atmosphere. The mixture was heated to reflux overnight. It was allowed to cool and concentrated. The solid obtained was dissolved in CHCl₃ and some drops of MeOH. Saturated NaHCO₃ was added and the aqueous phase was extracted 3 times with CHCl₃. The combined organic phases were dried over Na₂SO₄ and concentrated to dryness. The crude product was purified by chromatography on
20 silica gel using hexane-EtOAc mixtures of increasing polarity as eluent, to afford 0.22 g of the desired compound (yield: 41%).

LC-MS (método 1): t_R = 5.56 min; m/z = 383.0 [M+H]⁺.

EXAMPLE 73

25 2-(2,2-Diethoxyethyl)-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridine

EXAMPLE 74

1-(2,2-Diethoxyethyl)-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridine

In a volumetric flask 4,6-bis(4-fluorophenyl)-5-(4-pyridyl)-1H-pyrazolo[3,4-
30 b]pyridine (0.20 g, 0.5 mmol, obtained in example 1), KOH (0.03 g, 0.5 mmol), 2-bromo-1,1-diethoxyethane (0.10 g, 0.5 mmol) and 1-methoxy-2-(2-methoxyethoxy)ethane (2 mL) were introduced under argon atmosphere. The mixture was heated to 100 °C and stirred at this temperature overnight. It was

allowed to cool and a mixture of H₂O-EtOAc was added. The phases were separated and the aqueous phase was extracted with EtOAc. The combined organic phases were dried over Na₂SO₄ and concentrated to dryness. The crude product obtained was purified by chromatography on silica gel using hexane-EtOAc mixtures of increasing polarity as eluent, to afford 139 mg of 2-(2,2-diethoxyethyl)-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine (yield: 53%) and 60 mg of 1-(2,2-diethoxyethyl)-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine (yield: 24%).

Example 73: ¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.15 (t, J = 7.0 Hz, 6 H), 3.50 (m, 2 H), 3.73 (m, 2 H), 4.49 (d, J = 5.4 Hz, 2 H), 5.06 (t, J = 5.4 Hz, 1 H), 6.83 (dd, J_o = 1.4 Hz, J_m = 4.4 Hz, 2 H), 6.90 (t, J = 8.7 Hz, 2 H), 7.00 (t, J = 8.6 Hz, 2 H), 7.14 (m, 2 H), 7.29 (m, 2 H), 7.85 (s, 1 H), 8.32 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H).

Example 74: ¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.21 (t, J = 6.9 Hz, 6 H), 3.58 (m, 2 H), 3.82 (m, 2 H), 4.74 (d, J = 5.7 Hz, 2 H), 5.19 (t, J = 5.7 Hz, 1 H), 6.81 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H), 6.92 (t, J = 8.7 Hz, 2 H), 7.01 (t, J = 8.7 Hz, 2 H), 7.13 (m, 2 H), 7.25 (m, 2 H), 7.87 (s, 1 H), 8.32 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H).

EXAMPLE 75

4,6-Bis(4-fluorophenyl)-1-methyl-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine-3-carbonitrile

Following a similar procedure to that described in example 65, but using 3-bromo-4,6-bis(4-fluorophenyl)-1-methyl-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine (obtained in example 66) instead of 3-bromo-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)-1H-pyrazolo[3,4-*b*]pyridine, the title compound was obtained (yield: 48%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 4.32 (s, 3 H), 6.81 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H), 6.96 (t, J = 8.4 Hz, 2 H), 7.05 (t, J = 8.4 Hz, 2 H), 7.13 (m, 2 H), 7.29 (m, 2 H), 8.35 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H).

EXAMPLE 76

3-Bromo-6-(4-fluorophenyl)-5-(4-pyridyl)-1H-pyrazolo[3,4-*b*]pyridine

To a solution of 3-amino-6-(4-fluorophenyl)-5-(4-pyridyl)-1H-pyrazolo[3,4-*b*]pyridine (0.20 g, 0.7 mmol, obtained in example 70) in 48% HBr (1 mL) at 0 °C, a solution of NaNO₂ (0.05 g, 0.7 mmol) in water (0.1 mL) was added dropwise

over a period of 15 minutes maintaining the temperature at 0-5 °C. The mixture was stirred for 15 min at this temperature. Then, a solution of CuBr (0.24 g, 1.7 mmol) in 48% HBr (1 mL) was added slowly at 0°C. The resulting solution was stirred for 3 h at 0 °C. It was allowed to reach room temperature, neutralized at pH = 7 with saturated sodium bicarbonate and a 30% aqueous NH₃. It was filtered and the solid was washed with a mixture of CH₂Cl₂ and water. The organic phase was washed with 1 N NaOH and the aqueous phase extracted with EtOAc. The combined organic phases were dried over Na₂SO₄ and concentrated to dryness. The crude product obtained was purified by chromatography on silica gel using CHCl₃-MeOH mixtures of increasing polarity as eluent, to afford 50 mg of the title compound (yield: 21%).

¹H NMR (300 MHz, CDCl₃ + CD₃OD) δ (TMS): 4.24 (s, NH + CD₃OD), 6.92 (m, 2 H), 7.15 (m, 2 H), 7.29 (m, 2 H), 7.95 (s, 1 H), 8.39 (d, J = 6.0 Hz, 2 H).

EXAMPLE 77

15 6-Fluorophenyl-5-(4-pyridyl)-1H-pyrazolo[3,4-b]pyridine

3-Amino-6-(4-fluorophenyl)-5-(4-pyridyl)-1H-pyrazolo[3,4-b]pyridine (0.20 g, 0.7 mmol, obtained in example 70) and a solution of H₃PO₂ (0.2 mL, 2.0 mmol) in water (2 mL) were mixed and allowed to cool to 5 °C. A solution of NaNO₂ (0.10 g, 1.4 mmol) in water (0.4 mL) was added dropwise. This was stirred for 30 min at 5 °C, allowed to reach room temperature and stirred at room temperature for 4 h. It was neutralized with 1 N NaOH and extracted with CHCl₃. The combined organic phases were dried over Na₂SO₄ and concentrated to dryness. The crude product obtained was purified by chromatography on silica gel using increasing polarity hexane-EtOAc mixtures as eluent, to afford 63 mg of the title compound (yield: 33%).

¹H NMR (300 MHz, MeOH + CDCl₃) δ (TMS): 3.46 (s, NH + CD₃OD), 6.92 (t, J = 8.6 Hz, 2 H), 7.07 (dd, J_o = 1.6 Hz, J_m = 4.4 Hz, 2 H), 7.26 (m, 2 H), 8.08 (s, 1 H), 8.09 (s, 1 H), 8.37 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H).

EXAMPLE 78

30 N-Methyl-[3-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridin-2-yl]propyl]amine

To a solution of 2-(3-chloropropyl)-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridine (0.07 g, 0.1 mmol, obtained in example 16) in

acetonitrile (0.3 mL), methylamine (1.8 mL of a 33% solution in EtOH, 14.6 mmol) was added under argon atmosphere. This was heated to 60 °C for 3 days, adding methylamine (0.9 mL and 3.6 mL of a 33% solution in EtOH) after 24 and 48 h respectively. The organic phase was dried over Na₂SO₄ and concentrated to dryness. The crude product obtained was purified by silica gel EtOAc-MeOH mixtures of using increasing polarity as eluent, to afford 24 mg of the title compound in solid form (yield: 29%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 2.00 (broad s, NH + H₂O), 2.53 (m, 2 H), 2.61 (s, 3 H), 2.99 (t, J = 6.7 Hz, 2 H), 4.67 (t, J = 6.4 Hz, 2 H), 6.82 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H), 6.90 (t, J = 8.7 Hz, 2 H), 7.00 (t, J = 8.6 Hz, 2 H), 7.13 (m, 2 H), 7.27 (m, 2 H), 7.90 (s, 1 H), 8.32 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H).

EXAMPLE 79

[4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridin-2-yl]methanol

A suspension of 4,6-bis(4-fluorophenyl)-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine (0.20 g, 0.5 mmol, obtained in example 1) in 30-40% aqueous formaldehyde (0.9 mL) was stirred at 130 °C under argon atmosphere for 4h. The solvent was concentrated and the residue was dissolved in a mixture of CHCl₃ and water and the phases were separated. The organic phase was dried over Na₂SO₄ and concentrated to dryness. The crude product obtained was purified on chromatography on silica gel using hexane-EtOAc mixtures of increasing polarity as eluent, to afford 140 mg of the title compound in solid form (yield: 65%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 3.5 (t, 1 H, OH), 6.11 (d, J = 7.8 Hz, 2 H), 6.85 (dd, J_o = 1.8 Hz, J_m = 4.5 Hz, 2 H), 6.97 (t, J = 8.7 Hz, 1 H), 7.08 (t, J = 8.7 Hz, 1 H), 7.16 (m, 2 H), 7.31 (m, 2 H), 7.97 (s, 1 H), 8.37 (dd, J_o = 1.6 Hz, J_m = 4.3 Hz, 2 H).

EXAMPLE 80

2-[4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]-*N,N*-dimethylacetamide

EXAMPLE 81

2-[4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-1-yl]-*N,N*-dimethylacetamide

Following a similar procedure to that described in examples 73 and 74, but using 2-chloro-*N,N*-dimethylacetamide instead of 2-bromo-1,1-diethoxyethane, the title compounds were obtained.

Example 80: yield: 10%

- 5 Example 81: yield 32%; ¹H NMR (300 MHz, CDCl₃) δ (TMS): 3.01 (s, 3 H), 3.18 (s, 3 H), 5.47 (s, 2 H), 6.79 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H), 6.91 (t, J = 8.7 Hz, 1 H), 7.01 (t, J = 8.6 Hz, 1 H), 7.15 (m, 2 H), 7.23 (m, 2 H), 7.95 (s, 1 H), 8.31 (dd, J_o = 1.6 Hz, J_m = 4.3 Hz, 2 H).

EXAMPLE 82

- 10 4,6-Bis(4-fluorophenyl)-2-[2-(2-methoxyethoxy)ethyl]-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine

EXAMPLE 83

- 4,6-Bis(4-fluorophenyl)-1-[2-(2-methoxyethoxy)ethyl]-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine

- 15 Following a similar procedure to that described in examples 6 and 7, but using 1-bromo-2-(2-methoxyethoxy)ethane instead of iodoethane, the title compounds were obtained.

- Example 82: yield: 27%; ¹H NMR (300 MHz, CDCl₃) δ (TMS): 3.28 (s, 3 H), 3.45 (m, 2 H), 3.57 (m, 2 H), 4.05 (t, J = 5.1 Hz, 2 H), 4.63 (t, J = 5.1 Hz, 2 H), 6.82 (dd, 20 J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H), 6.89 (t, J = 8.7 Hz, 2 H), 6.99 (t, J = 8.7 Hz, 2 H), 7.14 (m, 2 H), 7.29 (m, 2 H), 7.93 (s, 1 H), 8.31 (dd, J_o = 1.6 Hz, J_m = 4.4 Hz, 2 H).
Example 83: yield: 19%; ¹H NMR (300 MHz, CDCl₃) δ (TMS): 3.32 (s, 3 H), 3.50 (m, 2 H), 3.68 (m, 2 H), 4.08 (t, J = 6.0 Hz, 2 H), 4.81 (t, J = 6.0 Hz, 2 H), 6.81 (dd, J_o = 1.4 Hz, J_m = 4.4 Hz, 2 H), 6.92 (t, J = 8.7 Hz, 2 H), 7.01 (t, J = 8.6 Hz, 2 H), 25 7.13 (m, 2 H), 7.26 (m, 2 H), 7.86 (s, 1 H), 8.32 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H).

EXAMPLE 84

- 4,6-Bis(4-fluorophenyl)-2-[3-(morpholin-4-yl)propyl]-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine

- a) 3-[4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]propyl
30 methanesulfonate

Following a similar procedure to that described in example 12 section b, but using 3-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]propan-1-ol

(obtained in example 18) instead of 2-[1-(*tert*-butoxycarbonyl)piperidin-4-yl]ethanol, the desired compound was obtained (yield: quantitative).

b) Title compound

In a volumetric flask, 3-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]propyl methanesulfonate (0.10 g, 0.2 mmol, obtained in section a), NaI (0.003 g, 0.02 mmol), morpholine (0.03 g, 0.4 mmol) and 1,2-dimethoxyethane (2 mL) were introduced under argon atmosphere. The mixture was heated to 90 °C and stirred at this temperature overnight. A mixture of water and EtOAc was added. The phases were separated. The aqueous phase was extracted with EtOAc. The organic phase was dried over Na₂SO₄ and concentrated to dryness. The crude product obtained was purified by chromatography on silica gel using increasing polarity mixtures of EtOAc-MeOH as eluent, to afford 37 mg of the title compound (yield: 36%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 2.27 (m, 2 H), 2.30 – 2.43 (complex signal, 6 H), 3.69 (m, 4 H), 4.53 (t, J = 6.6 Hz, 2 H), 6.83 (dd, J_o = 1.6 Hz, J_m = 4.4 Hz, 2 H), 6.89 (t, J = 8.7 Hz, 2 H), 7.00 (t, J = 8.6 Hz, 2 H), 7.14 (m, 2 H), 7.31 (m, 2 H), 7.81 (s, 1 H), 8.32 (dd, J_o = 1.6 Hz, J_m = 4.4 Hz, 2 H).

EXAMPLE 85

4,6-Bis(6-chloropyridin-3-yl)-2-methyl-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine

EXAMPLE 86

4,6-Bis(6-chloropyridin-3-yl)-1-methyl-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine

Following a similar procedure to that described in examples 73 and 74, but using 4,6-bis(6-chloropyridin-3-yl)-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine (obtained in example 40) instead of 4,6-bis(4-fluorophenyl)-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine and iodomethane instead of 2-bromo-1,1-diethoxyethane, the title compounds were obtained.

Example 85: yield: 26%; ¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.56 (s, NH + H₂O), 4.32 (s, 3 H), 6.88 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H), 7.19 (d, J = 8.4 Hz, 1 H), 7.31 (m, 2 H), 7.58 (dd, J_o = 2.4 Hz, J_m = 8.4 Hz, 1 H), 7.84 (s, 1 H), 8.37 (dd, J_o = 2.4 Hz, J_m = 6.9 Hz, 2 H), 8.43 (dd, J_o = 1.6 Hz, J_m = 4.4 Hz, 2 H).

Example 86: yield: 27%; ¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.57 (s, NH + H₂O), 4.26 (s, 3 H), 6.88 (dd, J_o = 1.8 Hz, J_m = 4.5 Hz, 2 H), 7.20 (d, J = 8.4 Hz, 1 H), 7.29 (d, J = 8.1 Hz, 1 H), 7.35 (dd, J_o = 2.4 Hz, J_m = 8.1 Hz, 1 H), 7.50 (dd, J_o =

2.6 Hz, $J_m = 8.2$ Hz, 1 H), 7.91 (s, 1 H), 8.34 (d, $J = 2.1$ Hz, 1 H), 8.43 – 8.45 (complex signal, 3 H)

EXAMPLE 87

4,6-Bis(6-chloropyridin-3-yl)-3-methyl-5-(4-pyridyl)-1H-pyrazolo[3,4- b]pyridine

5

Following a similar procedure to that described in example 1 method A, but using 1-(6-chloropyridin-3-yl)-2-(4-pyridyl)ethanone (obtained in example reference 9) instead of 1-(4-fluorophenyl)-2-(4-pyridyl)ethanone and 3-amino-5-methyl-2H-pyrazole instead of 3-amino-2H-pyrazole, the title compound was obtained (yield: 9%).

10

^1H NMR (300 MHz, $\text{CDCl}_3 + \text{CD}_3\text{OD}$) δ (TMS): 2.09 (s, 3 H), 3.51 (s, NH + H_2O), 6.92 (m, 2 H), 7.23 (d, $J = 8.4$ Hz, 1 H), 7.32 (m, 1 H), 7.51 (m, 1 H), 7.54 (dd, $J_o = 2.4$ Hz, $J_m = 8.4$ Hz, 1 H), 8.26 (broad s, 1 H), 8.35 – 8.38 (complex signal, 3 H).

EXAMPLE 88

4,6-Bis(6-methylpyridin-3-yl)-5-(4-pyridyl)-1H-pyrazolo[3,4-b]pyridine

15

Following a similar procedure to that described in example 1 method A, but using 1-(6-methylpyridin-3-yl)-2-(4-pyridyl)ethanone (obtained in reference example 13) instead of 1-(4-fluorophenyl)-2-(4-pyridyl)ethanone, the title compound was obtained (yield: 23%).

^1H NMR (300 MHz, CDCl_3) δ (TMS): 1.59 (s, NH + H_2O), 2.59 (s, 6 H), 6.90 (dd, $J_o = 1.5$ Hz, $J_m = 4.5$ Hz, 2 H), 6.99 (d, $J = 8.1$ Hz, 1 H), 7.09 (d, $J = 7.8$ Hz, 1 H), 7.33 (m, 2 H), 7.99 (s, 1 H), 8.39 (dd, $J_o = 1.5$ Hz, $J_m = 4.5$ Hz, 2 H), 8.47 (d, $J = 2.1$ Hz, 1 H), 8.80 (broad s, 1 H).

EXAMPLE 89

4,6-Bis(4-fluorophenyl)-2-(2-phthalimidoethyl)-5-(4-pyridyl)pyrazolo[3,4- b]pyridine

25

Following a similar procedure to that described in examples 6 and 7, but using *N*-(2-bromoethyl)phthalimide instead of iodomethane, the title compound was obtained (yield: 29%).

^1H NMR (300 MHz, CDCl_3) δ (TMS): 4.32 (t, $J = 6.1$ Hz, 2 H), 4.76 (t, $J = 6.0$ Hz, 2 H), 6.81 (dd, $J_o = 1.6$ Hz, $J_m = 4.4$ Hz, 2 H), 6.88 (t, $J = 8.7$ Hz, 2 H), 6.95 (t, $J = 8.6$ Hz, 2 H), 7.07 (m, 2 H), 7.25 (m, 2 H), 7.50 (m, 2 H), 7.61 (s, 1 H), 7.71 (m, 2 H), 8.32 (dd, $J_o = 1.6$ Hz, $J_m = 4.4$ Hz, 2 H).

30

EXAMPLE 90

2-(2-Aminoethyl)-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine

To a solution of 4,6-bis(4-fluorophenyl)-2-(2-phthalimidoethyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine (0.09 g, 0.2 mmol, obtained in example 89) in EtOH (2 mL), hydrazine monohydrate (0.02 g, 0.3 mmol) was added and the mixture was heated to reflux for 3 h. A mixture of water and EtOAc was added. The phases were separated. The aqueous phase was extracted with EtOAc. The organic phase was dried over Na₂SO₄ and concentrated to dryness. The crude product obtained was purified by chromatography on silica gel using CHCl₃-MeOH mixtures of increasing polarity as eluent, to afford 57 mg of the title compound (yield: 83%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.57 (s, NH₂ + H₂O), 3.38 (t, J = 5.7 Hz, 2 H), 4.49 (t, J = 5.5 Hz, 2 H), 6.83 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H), 6.90 (t, J = 8.8 Hz, 2 H), 7.00 (t, J = 8.7 Hz, 2 H), 7.15 (m, 2 H), 7.31 (m, 2 H), 7.85 (s, 1 H), 8.32 (dd, J_o = 1.6 Hz, J_m = 4.6 Hz, 2 H).

EXAMPLE 91

2-[4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethanol

A solution of 4,6-bis(4-fluorophenyl)-5-(4-pyridyl)-2-[2-(tetrahydropyran-2-yloxy)ethyl]pyrazolo[3,4-*b*]pyridine (1.08 g, 2.09 mmol, obtained in example 24) in a 4:2:1 mixture of AcOH:THF:H₂O (42 mL) was heated to 55 °C overnight. The mixture was allowed to cool and basified with saturated NaHCO₃ and extracted with EtOAc. The organic phase was dried over Na₂SO₄ and concentrated to dryness. The crude product obtained was purified by chromatography on silica gel using hexane-EtOAc mixtures of increasing polarity as eluent, to afford 0.78 g of the title compound (yield: 87%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.61 (broad s, OH + H₂O), 4.21 (t, J = 4.7 Hz, 2 H), 4.58 (t, J = 4.7 Hz, 2 H), 6.83 (dd, J_o = 1.6 Hz, J_m = 4.4 Hz, 2 H), 6.91 (t, J = 8.7 Hz, 2 H), 7.01 (t, J = 8.6 Hz, 2 H), 7.14 (m, 2 H), 7.27 (m, 2 H), 7.87 (s, 1 H), 8.33 (dd, J_o = 1.6 Hz, J_m = 4.5 Hz, 2 H).

EXAMPLE 92

6-(4-Fluorophenyl)-2-methyl-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine

Following a similar procedure to that described in examples 6 and 7, but using 6-fluorophenyl-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine (obtained in example 77)

instead of 4,6-bis(4-fluorophenyl)-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine and iodomethane instead of iodoethane, the title compound was obtained (yield: 35%).
¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.56 (broad s, NH + H₂O), 4.31 (s, 3 H), 6.95 (t, J = 8.7 Hz, 2 H), 7.11 (d, J = 6.0 Hz, 2 H), 7.41 (m, 2 H), 8.01 (s, 1 H), 8.06 (s, 1 H), 8.51 (d, J = 6.0 Hz, 2 H).

EXAMPLE 93

4,6-Bis(4-fluorophenyl)-2-(3-phthalimidopropyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine

Following a similar procedure to that described in examples 6 and 7, but using *N*-(3-bromopropyl)phthalimide instead of iodoethane, the title compound was obtained (yield: 31%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 2.54 (q, J = 6.4 Hz, 2 H), 3.79 (t, J = 6.2 Hz, 2 H), 4.50 (t, J = 6.6 Hz, 2 H), 6.82 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H), 6.89 (t, J = 8.7 Hz, 2 H), 7.02 (t, J = 8.7 Hz, 2 H), 7.17 (m, 2 H), 7.28 (m, 2 H), 7.71 (m, 2 H), 7.83 (m, 2 H), 7.93 (s, 1 H), 8.32 (dd, J_o = 1.6 Hz, J_m = 4.5 Hz, 2 H).

EXAMPLE 94

2-[4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]acetaldehyde

In a volumetric flask, 2-(2,2-diethoxyethyl)-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine (0.31 g, 0.6 mmol, obtained in example 73) and 1 N HCl (2.6 mL) were introduced. The mixture was heated to 100 °C for 1 hour. It was allowed to cool, adjusted to pH = 7 and extracted with EtOAc. The organic phase was dried over Na₂SO₄ and concentrated to dryness. The crude product obtained was purified by chromatography on silica gel using hexane-EtOAc mixtures of increasing polarity eluent, to afford 166 mg of the title compound (yield: 65%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 5.26 (s, 2 H), 6.83 (dd, J_o = 1.4 Hz, J_m = 4.4 Hz, 2 H), 6.91 (t, J = 8.5 Hz, 2 H), 7.00 (t, J = 8.8 Hz, 2 H), 7.15 (m, 2 H), 7.30 (m, 2 H), 7.88 (s, 1 H), 8.31 (m, 2 H), 9.85 (s, 1 H).

EXAMPLE 95

2-(3-Aminopropyl)-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine

Following a similar procedure to that described in example 90, but using 4,6-bis(4-fluorophenyl)-2-(3-phthalimidopropyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine

(obtained in example 93) instead of 4,6-bis(4-fluorophenyl)-2-(2-phthalimidoethyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine, the title compound was obtained (yield: 49%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.54 (s, NH₂ + H₂O), 2.19 (m, 2 H), 2.76 (t, J = 6.7 Hz, 2 H), 4.57 (t, J = 6.7 Hz, 2 H), 6.83 (dd, J_o = 1.6 Hz, J_m = 4.4 Hz, 2 H), 6.90 (t, J = 8.7 Hz, 2 H), 7.00 (t, J = 8.7 Hz, 2 H), 7.13 (m, 2 H), 7.30 (m, 2 H), 7.81 (s, 1 H), 8.32 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H).

EXAMPLE 96

***N*-[4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridin-3-ylmethyl]-1-(*tert*-butoxycarbonyl)piperidine-4-carboxamide**

Following a similar procedure to that described in example 48, but using 1-(*tert*-butoxycarbonyl)piperidine-4-carboxylic acid instead of 2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]acetic acid and 3-aminomethyl-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine (obtained in example 68) instead of morpholine, the title compound was obtained (yield: 26%).

EXAMPLE 97

***N*-[4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridin-3-ylmethyl]piperidine-4-carboxamide**

Following a similar procedure to that described in example 36, but using *N*-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridin-3-ylmethyl]-1-(*tert*-butoxycarbonyl)piperidine-4-carboxamide (obtained in example 96) instead of 2-[1-(*tert*-butoxycarbonyl)piperidin-4-yl]-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine, the title compound was obtained (yield: 63%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.40 – 1.90 (complex signal, 3 H + H₂O), 2.18 (m, 2 H), 2.63 (m, 2 H), 3.13 (m, 2 H), 4.15 (d, J = 5.1 Hz, 2 H), 5.20 (broad s, 1H, NH), 6.20 (broad s, 1 H, NH), 6.78 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H), 6.94 (t, J = 8.7 Hz, 2 H), 7.03 (t, J = 8.7 Hz, 2 H), 7.08 – 7.14 (complex signal, 4 H), 8.30 (dd, J_o = 1.6 Hz, J_m = 4.4 Hz, 2 H).

EXAMPLE 98

2-(3-Benzoyloxypropyl)-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine

EXAMPLE 99

1-(3-Benzoyloxypropyl)-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine

Following a similar procedure to that described in examples 6 and 7, but using 3-bromopropanol benzylic ether instead of iodoethane, the title compounds were obtained.

Example 98: yield: 43%; ¹H NMR (300 MHz, CDCl₃) δ (TMS): 2.38 (m, 2 H), 3.48 (t, J = 5.6 Hz, 2 H), 4.44 (s, 2 H), 4.57 (t, J = 6.7 Hz, 2 H), 6.82 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H), 6.89 (t, J = 8.7 Hz, 2 H), 6.99 (t, J = 8.7 Hz, 2 H), 7.10 (m, 2 H), 7.28 – 7.32 (complex signal, 7 H), 7.73 (s, 1 H), 8.32 (dd, J_o = 1.8 Hz, J_m = 4.5 Hz, 2 H).

Example 99: yield: 20%; ¹H NMR (300 MHz, CDCl₃) δ (TMS): 2.19 (m, 2 H), 3.58 (t, J = 6.1 Hz, 2 H), 4.48 (s, 2 H), 4.75 (t, J = 6.9 Hz, 2 H), 6.81 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H), 6.90 (t, J = 8.7 Hz, 2 H), 7.01 (t, J = 8.6 Hz, 2 H), 7.12 (m, 2 H), 7.24 – 7.33 (complex signal, 7 H), 7.86 (s, 1 H), 8.32 (dd, J_o = 1.6 Hz, J_m = 4.4 Hz, 2 H).

EXAMPLE 100

15 *N,N*-Diethyl-[2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl]amine

EXAMPLE 101

N,N-Diethyl-[2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-1-yl]ethyl]amine

20 Following a similar procedure to that described in examples 6 and 7, but using *N*-(2-chloroethyl)diethylamine hydrochloride instead of iodoethane and 2 equivalents of KOH, the title compounds were obtained.

Example 100: yield: 5%; ¹H NMR (300 MHz, CDCl₃) δ (TMS): 0.99 (t, J = 7.2 Hz, 6 H), 2.56 (c, J = 7.1 Hz, 4 H), 3.08 (t, J = 6.5 Hz, 2 H), 4.48 (t, J = 6.5 Hz, 2 H), 6.82 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H), 6.89 (t, J = 8.8 Hz, 2 H), 7.00 (t, J = 8.6 Hz, 2 H), 7.14 (m, 2 H), 7.29 (m, 2 H), 7.85 (s, 1 H), 8.32 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H).

Example 101: yield: 73%; ¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.05 (t, J = 7.2 Hz, 6 H), 2.66 (c, J = 7.2 Hz, 4 H), 3.09 (t, J = 7.2 Hz, 2 H), 4.69 (t, J = 7.2 Hz, 2 H), 6.82 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H), 6.92 (t, J = 8.7 Hz, 2 H), 6.98 (t, J = 8.7 Hz, 2 H), 7.14 (m, 2 H), 7.26 (m, 2 H), 7.85 (s, 1 H), 8.32 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H).

EXAMPLE 102

**4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)-2-(3-pyridylmethyl)pyrazolo[3,4-
b]pyridine**

EXAMPLE 103

**4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)-1-(3-pyridylmethyl)pyrazolo[3,4-
b]pyridine**

5

Following a similar procedure to that described in examples 6 and 7, but using 3-chloromethylpyridine hydrochloride instead of iodoethane, the title compounds were obtained.

Example 102: yield: 16%; ¹H NMR (300 MHz, CDCl₃) δ (TMS): 5.63 (s, 2 H), 6.81
10 (dd, J_o = 1.8 Hz, J_m = 4.5 Hz, 2 H), 6.90 (t, J = 8.8 Hz, 2 H), 6.99 (t, J = 8.7 Hz, 2 H), 7.10 (m, 2 H), 7.31 (m, 2 H), 7.82 (m, 2 H), 8.32 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H), 8.61 (dd, J_o = 1.5 Hz, J_m = 4.8 Hz, 2 H), 8.68 (s, 1 H).

Example 103: yield: 22%; ¹H NMR (300 MHz, CDCl₃) δ (TMS): 5.80 (s, 2 H), 6.81
15 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H), 6.94 (t, J = 8.6 Hz, 2 H), 7.01 (t, J = 8.7 Hz, 2 H), 7.13 (m, 2 H), 7.27 (m, 2 H), 7.79 (m, 1 H), 7.88 (s, 1 H), 8.32 (dd, J_o = 1.4 Hz, J_m = 4.4 Hz, 2 H), 8.56 (d, J = 5.0 Hz, 2 H), 8.75 (s, 1 H).

EXAMPLE 104

***N,N*-Dimethyl-[3-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-
2-yl]propyl]amine**

20

EXAMPLE 105

***N,N*-Dimethyl-[3-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-
1-yl]propyl]amine**

Following a similar procedure to that described in examples 6 and 7, but using *N*-(3-chloropropyl)dimethylamine hydrochloride instead of iodoethane, the title
25 compounds were obtained.

Example 104: 22%; ¹H NMR (300 MHz, CDCl₃) δ (TMS): 2.21 – 2.29 (complex
signal, 10 H), 4.51 (t, J = 6.6 Hz, 2 H), 6.82 (dd, J_o = 1.6 Hz, J_m = 4.4 Hz, 2 H),
6.89 (t, J = 8.8 Hz, 2 H), 7.00 (t, J = 8.7 Hz, 2 H), 7.14 (m, 2 H), 7.30 (m, 2 H),
7.81 (s, 1 H), 8.31 (dd, J_o = 1.6 Hz, J_m = 4.4 Hz, 2 H).

30 Example 105: yield: 19%; ¹H NMR (300 MHz, CDCl₃) δ (TMS): 2.18 – 2.27
(complex signal, 8 H), 2.43 (t, J = 7.2 Hz, 2 H), 4.65 (t, J = 7.1 Hz, 2 H), 6.82 (dd,
J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H), 6.93 (t, J = 8.8 Hz, 2 H), 7.01 (t, J = 8.7 Hz, 2 H),
7.14 (m, 2 H), 7.27 (m, 2 H), 7.86 (s, 1 H), 8.32 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H).

EXAMPLE 106**1-[2-[4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridin-2-yl]ethy]piperidin-4-ol**

In a volumetric flask, 2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridin-2-yl]acetaldehyde (0.08 g, 0.2 mmol, obtained in example 94), sodium triacetoxyborohydride (0.08 g, 0.4 mmol), 4-hydroxypiperidine (0.02 g, 0.2 mmol) and 1,2-dichloroethane (3 mL) were introduced under argon atmosphere. The mixture was stirred overnight at room temperature. It was concentrated and a mixture of water and EtOAc were added. The phases were separated. The aqueous phase was extracted with EtOAc. The organic phase was dried over Na₂SO₄ and concentrated to dryness. The crude product obtained was purified by chromatography on silica gel using increasing polarity mixtures of EtOAc-MeOH as eluent, to afford 19 mg of the title compound (yield: 20%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.57 (broad s, 1 H + OH + H₂O), 1.85 (m, 2 H), 2.05 (m, 2 H), 2.30 (m, 2 H), 2.80 (m, 2 H), 3.01 (m, 2 H), 4.54 (m, 2 H), 6.83 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H), 6.90 (t, J = 8.6 Hz, 2 H), 7.01 (t, J = 8.6 Hz, 2 H), 7.14 (m, 2 H), 7.29 (m, 2 H), 7.85 (s, 1 H), 8.34 (dd, J_o = 1.6 Hz, J_m = 4.4 Hz, 2 H).

EXAMPLE 107**3-[4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridin-2-yl]-2-hydroxypropan-1-ol****EXAMPLE 108****3-[4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridin-1-yl]-2-hydroxypropan-1-ol**

Following a similar procedure to that described in examples 6 and 7, but using 3-bromopropane-1,2-diol instead of iodoethane, the title compounds were obtained.

Example 107: yield: 17%; ¹H NMR (300 MHz, CDCl₃) δ (TMS): 3.49 (s, 1H, OH), 3.72 (m, 2 H), 4.32 (m, 1 H), 4.60 (m, 2 H), 5.30 (s, 1H, OH), 6.83 (dd, J_o = 1.6 Hz, J_m = 4.4 Hz, 2 H), 6.91 (t, J = 8.7 Hz, 2 H), 7.01 (t, J = 8.6 Hz, 2 H), 7.14 (m, 2 H), 7.28 (m, 2 H), 7.88 (s, 1 H), 8.32 (dd, J_o = 1.6 Hz, J_m = 4.4 Hz, 2 H).

Example 108: yield: 26%; ¹H NMR (300 MHz, CDCl₃) δ (TMS): 3.65 (m, 2 H), 4.26 (m, 1 H), 4.78 (m, 2 H), 5.30 (s, 2 H, OH), 6.82 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H),

6.94 (t, J = 8.6 Hz, 2 H), 7.03 (t, J = 8.7 Hz, 2 H), 7.14 (m, 2 H), 7.28 (m, 2 H), 7.91 (s, 1 H), 8.34 (dd, $J_o = 1.5$ Hz, $J_m = 4.5$ Hz, 2 H).

EXAMPLE 109

**4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)-2-(4-pyridylmethyl)pyrazolo[3,4-
b]pyridine**

EXAMPLE 110

**4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)-1-(4-pyridylmethyl)pyrazolo[3,4-
b]pyridine**

Following a similar procedure to that described in examples 6 and 7, but using 4-chloromethylpyridine hydrochloride instead of iodoethane, the title compounds were obtained.

Example 109: yield: 29%; ^1H NMR (300 MHz, CDCl_3) δ (TMS): 5.63 (s, 2 H), 6.83 (dd, $J_o = 1.5$ Hz, $J_m = 4.5$ Hz, 2 H), 6.90 (t, J = 8.7 Hz, 2 H), 6.99 (t, J = 8.7 Hz, 2 H), 7.12 (m, 2 H), 7.23 – 7.33 (complex signal, 4 H), 7.84 (s, 1 H), 8.32 (dd, $J_o = 1.5$ Hz, $J_m = 4.5$ Hz, 2 H), 8.61 (dd, $J_o = 1.5$ Hz, $J_m = 4.8$ Hz, 2 H).

Example 110: yield: 15%; 5.80 (s, 2 H), 6.82 (dd, $J_o = 1.5$ Hz, $J_m = 4.5$ Hz, 2 H), 6.93 (t, J = 8.7 Hz, 2 H), 7.00 (t, J = 8.7 Hz, 2 H), 7.14 (m, 2 H), 7.20 – 7.26 (complex signal, 4 H), 7.93 (s, 1 H), 8.34 (dd, $J_o = 1.5$ Hz, $J_m = 4.5$ Hz, 2 H), 8.61 (dd, $J_o = 1.5$ Hz, $J_m = 4.8$ Hz, 2 H).

EXAMPLE 111

***N*-(*tert*-Butoxycarbonyl)-[1-[3-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]propyl]piperidin-4-yl]amine**

To a solution of 3-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]propyl methanesulfonate (0.15 g, 0.3 mmol, obtained in example 84 section a) in acetonitrile (2 mL), *N*-(*tert*-butoxycarbonyl)-*N*-(4-piperidyl)amine (0.12 g, 0.6 mmol) was added under argon atmosphere and heated to 60 °C overnight. A mixture of CHCl_3 and saturated NaHCO_3 were added. The phases were separated. The aqueous phase was extracted with CHCl_3 . The organic phase was dried over Na_2SO_4 and concentrated to dryness. The crude product obtained was purified by chromatography on silica gel using increasing polarity mixtures of EtOAc-MeOH as eluent, to afford 40 mg of the title compound (yield: 22%).

^1H NMR (300 MHz, CDCl_3) δ (TMS): 1.44 (broad s, 2 H + NH_2 + H_2O), 1.95 (m, 2 H), 2.06 (m, 2 H), 2.25 (m, 2 H), 2.35 (m, 2 H), 3.46 (m, 2 H), 4.40 (m, 1 H), 4.51

(t, 2 H), 6.83 (d, J = 6.0 Hz, 2 H), 6.90 (t, J = 8.7 Hz, 2 H), 7.00 (t, J = 8.7 Hz, 2 H), 7.14 (m, 2 H), 7.29 (m, 2 H), 7.80 (s, 1 H), 8.32 (d, J = 6.0 Hz, 2 H).

EXAMPLE 112

2-[1-(*tert*-Butoxycarbonyl)piperidin-4-yl]-6-(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine

EXAMPLE 113

1-[1-(*tert*-Butoxycarbonyl)piperidin-4-yl]-6-(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine

Following a similar procedure to that described in examples 6 and 7, but using 6-fluorophenyl-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine (obtained in example 77) instead of 4,6-bis(4-fluorophenyl)-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine and 1-(*tert*-butoxycarbonyl)piperidin-4-yl methanesulfonate (obtained in example 14 section a) instead of iodoethane, the title compounds were obtained.

Example 112: yield: 26%; ¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.50 (s, 9 H), 2.17 – 2.32 (complex signal, 2 H), 2.99 (m, 2 H), 4.33 (m, 2 H), 4.60 (m, 1 H), 6.95 (t, J = 8.7 Hz, 2 H), 7.11 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H), 7.41 (m, 2 H), 8.06 (s, 1 H), 8.07 (s, 1 H), 8.52 (dd, J_o = 1.6 Hz, J_m = 4.4 Hz, 2 H).

Example 113: yield: 55%; ¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.50 (s, 9 H), 2.04 (m, 2 H), 2.30 (m, 2 H), 3.00 (m, 2 H), 4.42 (m, 2 H), 5.10 (m, 1 H), 7.00 (t, J = 8.6 Hz, 2 H), 7.10 (dd, J_o = 1.6 Hz, J_m = 4.5 Hz, 2 H), 7.37 (m, 2 H), 8.07 (s, 1 H), 8.10 (s, 1 H), 8.52 (dd, J_o = 1.8 Hz, J_m = 4.5 Hz, 2 H).

EXAMPLE 114

3-Methyl-4,6-bis(6-methylpyridin-3-yl)-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine

Following a similar procedure to that described in example 1 method A, but using 1-(6-methylpyridin-3-yl)-2-(4-pyridyl)ethanone (obtained in reference example 13) instead of 1-(4-fluorophenyl)-2-(4-pyridyl)ethanone and 3-amino-5-methyl-2*H*-pyrazole instead of 3-amino-2*H*-pyrazole, the title compound was obtained (yield: 24%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 2.07 (s, 3 H), 2.54 (s, 3 H), 2.57 (s, 3 H), 6.85 (m, 2 H), 6.99 (d, J = 8.1 Hz, 1 H), 7.08 (d, J = 8.1 Hz, 1 H), 7.30 (m, 1 H), 8.10 (dd, J_o = 2.4 Hz, J_m = 8.1 Hz, 1 H), 8.32 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H), 8.38 (s, 1 H), 8.57 (s, 1 H), 10.74 (broad s, NH).

EXAMPLE 115**1-[3-[4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]propyl]piperidin-4-one**

In a volumetric flask, 3-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]propyl methanesulfonate (0.15 g, 0.3 mmol, obtained in example 84 section a), NaI (0.008 g, 0.06 mmol), 4-piperidone monohydrate hydrochloride (0.04 g, 0.3 mmol), K₂CO₃ (0.07 g, 0.5 mmol) and DMF (2 mL) were introduced under argon atmosphere. The mixture was heated to 60 °C for 24 h. It was allowed to cool and a mixture of water and EtOAc was added. The phases were separated. The aqueous phase was extracted with EtOAc. The organic phase was dried over Na₂SO₄ and concentrated to dryness. The crude product obtained was purified by chromatography on silica gel using EtOAc-MeOH mixtures of increasing polarity as eluent, to afford 16 mg of the title compound (yield: 10%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.57 (broad s, 4 H + H₂O), 2.30 (m, 2 H), 2.42 (m, 2 H), 2.48 (m, 2 H), 2.73 (t, 2 H), 4.55 (t, 2 H), 6.83 (d, J = 6.0 Hz, 2 H), 6.90 (t, J = 8.7 Hz, 2 H), 7.00 (t, J = 8.6 Hz, 2 H), 7.18 (m, 2 H), 7.27 (m, 2 H), 7.82 (s, 1 H), 8.32 (d, J = 6.0 Hz, 2 H).

EXAMPLE 116***N*-(*tert*-Butoxycarbonyl)-[1-[2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl]piperidin-4-yl]amine****a) 2-[4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl methanesulfonate**

Following a similar procedure to that described in example 12 section b, but using 2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethanol (obtained in example 91) instead of 2-[1-(*tert*-butoxycarbonyl)piperidin-4-yl]ethanol, the desired compound was obtained (yield: quantitative).

b) Title compound

Following a similar procedure to that described in example 111, but using 2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl methanesulfonate (obtained in section a) instead of 3-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]propyl methanesulfonate the desired compound was obtained (yield: 66%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.44 (s, 9 H), 1.57 (broad s, 1 H + NH + H₂O), 1.90 (m, 2 H), 2.22 (m, 2 H), 2.82 (m, 2 H), 3.01 (t, J = 6.6 Hz, 2 H), 3.45 (m, 1 H), 4.40 (m, 1 H), 4.52 (t, J = 6.5 Hz, 2 H), 6.82 (d, J = 5.7 Hz, 2 H), 6.89 (t, J = 8.7 Hz, 2 H), 7.01 (t, J = 8.6 Hz, 2 H), 7.14 (m, 2 H), 7.29 (m, 2 H), 7.82 (s, 1 H),
5 8.31 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H).

EXAMPLE 117

***N*-Methyl-[1-[2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl]piperidin-4-yl]amine**

Following a similar procedure to that described in example 111, but using 2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl methanesulfonate
10 (obtained in example 116 section a) instead of 3-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]propyl methanesulfonate and *N*-methyl-*N*-(piperidin-4-yl)amine instead of *N*-(*tert*-butoxycarbonyl)-*N*-(4-piperidyl)amine, the desired compound was obtained (yield: 50%).

15 ¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.61 (broad s, 1 H + NH + H₂O), 2.28 (s, 3 H), 2.41 (m, 4 H), 2.57 (m, 4 H), 3.03 (t, J = 6.5 Hz, 2 H), 4.55 (t, J = 6.5 Hz, 2 H), 6.83 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H), 6.90 (t, J = 8.7 Hz, 2 H), 7.01 (t, J = 8.7 Hz, 2 H), 7.14 (m, 2 H), 7.29 (m, 2 H), 7.85 (s, 1 H), 8.32 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H).

20

EXAMPLE 118

[1-[3-[4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]propyl]piperidin-4-yl]amine

Following a similar procedure to that described in example 36, but using *N*-(*tert*-butoxycarbonyl)-[1-[3-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]propyl]piperidin-4-yl]amine (obtained in example 111) instead of 2-[1-(*tert*-
25 butoxycarbonyl)piperidin-4-yl]-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine, the title compound was obtained (yield: 89%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.35 (m, 2 H), 1.62 (broad s NH₂ + H₂O), 1.85 (m, 2 H), 2.03 (m, 2 H), 2.25 (m, 2 H), 2.36 (m, 2 H), 2.70 (m, 1 H), 2.82 (m, 2 H),
30 4.52 (d, J = 6.6 Hz, 2 H), 6.83 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H), 6.90 (t, J = 8.7 Hz, 2 H), 7.00 (t, J = 8.6 Hz, 2 H), 7.13 (m, 2 H), 7.29 (m, 2 H), 7.81 (s, 1 H), 8.31 (dd, J_o = 1.6 Hz, J_m = 4.4 Hz, 2 H).

EXAMPLE 119

2-[1-[2-[4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl]piperidin-4-yl]ethanol

Following a similar procedure to that described in example 111, but using 2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl methanesulfonate (obtained in example 116 section a) instead of 3-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]propyl methanesulfonate and 2-(4-piperidyl)ethanol instead of *N*-(*tert*-butoxycarbonyl)-*N*-(4-piperidyl)amine, the desired compound was obtained (yield: 50%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.48 – 1.53 (complex signal, 5 H), 2.11 (m, 2 H), 2.88 (m, 2 H), 2.99 (m, 2 H), 3.70 (t, *J* = 6.5 Hz, 2 H), 4.55 (t, *J* = 6.5 Hz, 2 H), 6.83 (dd, *J*_o = 1.5 Hz, *J*_m = 4.5 Hz, 2 H), 6.89 (t, *J* = 8.7 Hz, 2 H), 7.00 (t, *J* = 8.6 Hz, 2 H), 7.14 (m, 2 H), 7.29 (m, 2 H), 7.86 (s, 1 H), 8.32 (dd, *J*_o = 1.5 Hz, *J*_m = 4.5 Hz, 2 H).

EXAMPLE 120

[1-[2-[4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl]piperidin-4-yl]amine

Following a similar procedure to that described in example 36, but using *N*-(*tert*-butoxycarbonyl)-[1-[2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl]piperidin-4-yl]amine (obtained in example 116) instead of 2-[1-(*tert*-butoxycarbonyl)piperidin-4-yl]-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine, the title compound was obtained (yield: 77%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.38 (m, 2 H), 1.78 (broad s, 2 H + NH₂ + H₂O), 2.18 (m, 2 H), 2.75 (m, 1 H), 2.84 (m, 2 H), 3.02 (t, *J* = 6.3 Hz, 2 H), 4.54 (t, *J* = 6.5 Hz, 2 H), 6.82 (d, *J* = 5.7 Hz, 2 H), 6.89 (t, *J* = 8.7 Hz, 2 H), 7.00 (t, *J* = 8.6 Hz, 2 H), 7.14 (m, 2 H), 7.29 (m, 2 H), 7.84 (s, 1 H), 8.31 (d, *J* = 6.0 Hz, 2 H).

EXAMPLE 121

6-(4-Fluorophenyl)-2-(4-piperidyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine

Following a similar procedure to that described in example 36, but using 2-[1-(*tert*-butoxycarbonyl)piperidin-4-yl]-6-(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine (obtained in example 112) instead of 2-[1-(*tert*-butoxycarbonyl)piperidin-4-yl]-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine, the title compound was obtained (yield: 77%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.89 (m, NH + H₂O), 2.25 (m, 2 H), 2.30 (m, 2 H), 2.89 (m, 2 H), 3.34 (m, 2 H), 4.60 (m, 1 H), 6.95 (t, J = 8.7 Hz, 2 H), 7.11 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H), 7.40 (m, 2 H), 8.07 (s, 1 H), 8.08 (s, 1 H), 8.51 (dd, J_o = 1.6 Hz, J_m = 4.4 Hz, 2 H).

5

EXAMPLE 122**6-(4-Fluorophenyl)-1-(4-piperidyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine**

Following a similar procedure to that described in example 36, but using 1-[1-(*tert*-butoxycarbonyl)piperidin-4-yl]-6-(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-

10 4-yl]-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine, the title compound was obtained (yield: quantitative).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.60 (m, NH + H₂O), 2.42 (m, 2 H), 2.67 (m, 2 H), 3.28 (m, 2 H), 3.72 (m, 2 H), 5.28 (m, 1 H), 7.00 (t, J = 8.7 Hz, 2 H), 7.11 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H), 7.37 (m, 2 H), 8.10 (s, 1 H), 8.12 (s, 1 H), 8.54 (dd, 15 J_o = 1.6 Hz, J_m = 4.4 Hz, 2 H).

EXAMPLE 123**3-Amino-5-[2-(methylsulfanyl)pyrimidin-4-yl]-6-[3-(trifluoromethyl)phenyl]-1*H*-pyrazolo[3,4-*b*]pyridine**

Following a similar procedure to that described in reference example 15 section c, 20 but using 2-chloro-5-[2-(methylsulfanyl)pyrimidin-4-yl]-6-[3-(trifluoromethyl)phenyl]pyridine-3-carbonitrile (obtained in reference example 14) instead of 3-(1-benzylpiperidin-4-yl)-3-oxopropiononitrile, the title compound was obtained (yield: 78%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 2.44 (s, 3 H), 4.32 (broad s, NH₂), 6.61 (d, J = 25 5.1 Hz, 1 H), 7.44 – 7.84 (complex signal, 2 H), 7.65 (d, J = 7.2 Hz, 1 H), 7.84 (s, 1 H), 8.30 (d, J = 5.1 Hz, 1 H), 8.40 (s, 1 H), 9.55 (broad s, NH).

EXAMPLE 124**4,6-Bis(4-fluorophenyl)-2-[3-[1-(*tert*-butoxycarbonyl)piperazin-4-yl]propyl]-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine**

30 Following a similar procedure to that described in example 84 section b, but using 1-(*tert*-butoxycarbonyl)piperazine instead of morpholine and adding triethylamine (1.5 equivalents), the desired compound was obtained (yield: 14%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 2.66 (m, 2 H), 2.98 (m, 4 H), 3.10 (m, 2 H), 3.66 – 3.74 (complex signal, 4 H), 4.68 (m, 2 H), 6.85 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H), 6.91 (t, J = 8.7 Hz, 2 H), 7.00 (t, J = 8.7 Hz, 2 H), 7.14 (m, 2 H), 7.28 (m, 2 H), 7.98 (s, 1 H), 8.32 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H).

5

EXAMPLE 125**4,6-Bis(4-fluorophenyl)-2-[3-(piperazin-1-yl)propyl]-5-(4-pyridyl)pyrazolo[3,4-b]pyridine**

Following a similar procedure to that described in example 36, but using 4,6-bis(4-fluorophenyl)-2-[3-[1-(*tert*-butoxycarbonyl)piperazin-4-yl]propyl]-5-(4-

10

pyridyl)pyrazolo[3,4-*b*]pyridine (obtained in example 124) instead of 2-[1-(*tert*-butoxycarbonyl)piperidin-4-yl]-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine, the title compound was obtained (yield: quantitative).

15

¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.78 (broad s, 2 H + NH + H₂O), 2.28 (m, 2 H), 2.40 – 2.59 (complex signal, 6 H), 3.04 (m, 2 H), 4.52 (m, 2 H), 6.83 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H), 6.90 (t, J = 8.7 Hz, 2 H), 7.00 (t, J = 8.7 Hz, 2 H), 7.14 (m, 2 H), 7.29 (m, 2 H), 7.80 (s, 1 H), 8.31 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H).

EXAMPLE 126**5-[2-(Methylsulfanyl)pyrimidin-4-yl]-6-[3-(trifluoromethyl)phenyl]-1H-pyrazolo[3,4-*b*]pyridine**

20

Following a similar procedure to that described in example 77, but using 3-amino-5-[2-(methylsulfanyl)pyrimidin-4-yl]-6-[3-(trifluoromethyl)phenyl]-1H-pyrazolo[3,4-*b*]pyridine (obtained in example 123) instead of 3-amino-6-(4-fluorophenyl)-5-(4-pyridyl)-1H-pyrazolo[3,4-*b*]pyridine, the title compound was obtained (yield: 34%).

25

¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.55 (broad s, NH + H₂O), 6.70 (d, J = 5.1 Hz, 1 H), 7.50 (m, 2 H), 7.72 (m, 1 H), 7.86 (s, 1 H), 8.24 (s, 1 H), 8.35 (d, J = 5.1 Hz, 1 H), 8.54 (s, 1 H).

EXAMPLE 127**5-[2-(Methylsulfonyl)pyrimidin-4-yl]-6-[3-(trifluoromethyl)phenyl]-1H-pyrazolo[3,4-*b*]pyridine**

30

Following a similar procedure to that described in example 56, but using 5-[2-(methylsulfanyl)pyrimidin-4-yl]-6-[3-(trifluoromethyl)phenyl]-1H-pyrazolo[3,4-*b*]pyridine (obtained in example 126) instead of 4,6-bis(4-fluorophenyl)-2-(4-

methylsulfanylphenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine and 2 equivalents of *m*-chloroperbenzoic acid, the title compound was obtained (yield: quantitative).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.56 (broad s, NH + H₂O), 3.22 (s, 3 H), 7.21 (d, J = 5.1 Hz, 1 H), 7.49 (d, J = 4.8 Hz, 2 H), 7.71 (m, 1 H), 7.84 (s, 1 H), 8.30 (s, 1 H), 8.71 (m, 2 H).

EXAMPLE 128

(1*S*)-*N*-(1-Phenylethyl)-[4-[6-[3-(trifluoromethyl)phenyl]-1*H*-pyrazolo[3,4-*b*]pyridin-5-yl]pyrimidin-2-yl]amine

A mixture of 5-[2-(methylsulfonyl)pyrimidin-4-yl]-6-[3-(trifluoromethyl)phenyl]-1*H*-pyrazolo[3,4-*b*]pyridine (0.55 g, 0.13 mmol, obtained in example 127) and (1*S*)-1-phenylethylamine (0.16 g, 1.3 mmol) was heated to 100 °C for 1 h. It was allowed to cool and the crude product obtained was purified by chromatography on silica gel using increasing polarity mixtures of EtOAc-hexane as eluent, to afford 10 mg of the title compound (yield: 16%).

¹H NMR (300 MHz, CDCl₃ + CD₃OD) δ (TMS): 1.50 (d, 3 H), 3.80 (broad s, 2 NH + H₂O), 6.30 (d, 1 H), 7.20 – 7.40 (complex signal, 6 H), 7.50 (d, 1 H), 7.68 (d, 1 H), 7.76 (d, 1 H), 7.83 (s, 1 H), 8.08 (d, 1 H), 8.15 (s, 1 H), 8.24 (broad s, 1 H).

EXAMPLE 129

1-[3-[4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]propyl]piperidin-4-ol

Following a similar procedure to that described in example 111, but using piperidin-4-ol instead of *N*-(*tert*-butoxycarbonyl)-*N*-(4-piperidyl)amine, the desired compound was obtained (yield: 39%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.57 (m, 2 H), 1.88 – 2.50 (complex signal, 5 H + OH + H₂O), 2.75 (m, 2 H), 3.44 – 3.51 (complex signal, 4 H), 4.52 (t, J = 6.6 Hz, 2 H), 6.83 (dd, J_o = 1.6 Hz, J_m = 4.4 Hz, 2 H), 6.89 (t, J = 8.7 Hz, 2 H), 7.00 (t, J = 8.6 Hz, 2 H), 7.14 (m, 2 H), 7.34 (m, 2 H), 7.82 (s, 1 H), 8.32 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H).

EXAMPLE 130

2-[1-[3-[4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]propyl]piperidin-4-yl]ethanol

Following a similar procedure to that described in example 111, but using 2-(4-piperidyl)ethanol instead of *N*-(*tert*-butoxycarbonyl)-*N*-(4-piperidyl)amine, the desired compound was obtained (yield: 44%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.20-2.00 (broad s, 8 H + OH + H₂O), 2.50 (m, 1 H), 2.85 (m, 1 H), 3.50 (m, 1 H), 3.70 (m, 4 H), 4.59 (m, 2 H), 6.83 (d, J = 6.0 Hz, 2 H), 6.90 (t, J = 8.7 Hz, 2 H), 7.00 (t, J = 8.6 Hz, 2 H), 7.14 (m, 2 H), 7.30 (m, 2 H), 7.89 (s, 1 H), 8.32 (d, J = 6.0 Hz, 2 H).

EXAMPLE 131

4,6-Bis(4-fluorophenyl)-3-(4-piperidyl)-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine

a) 3-(1-Benzylpiperidin-4-yl)-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine

Following a similar procedure to that described in example 1 method A, but using 3-amino-5-(1-benzylpiperidin-4-yl)-2*H*-pyrazole (obtained in reference example 15)

Instead of 3-amino-2*H*-pyrazole, the title compound was obtained (yield: 6%).

b) Title compound

To a solution of 3-(1-benzylpiperidin-4-yl)-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine (23 mg, 0.04 mmol, obtained in the previous section) in MeOH (1 mL), Pd/C and a solution of HCOONH₄ (0.01 g, 0.2 mmol) in water (0.06 mL) were added under argon atmosphere. The mixture was heated to reflux for 5h. It was filtered through celite and concentrated. The residue was dissolved in CHCl₃ and washed with saturated NaHCO₃, to afford 2 mg of the title compound (yield: 10%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.10 – 1.90 (broad s, 7 H), 2.23 (m, 2 H), 2.99 (m, 2 H), 6.80 (dd, J_o = 1.6 Hz, J_m = 4.4 Hz, 2 H), 6.92 (t, J = 8.7 Hz, 2 H), 7.02 (t, J = 8.7 Hz, 2 H), 7.12 (m, 2 H), 7.28 (m, 2 H), 8.28 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H).

EXAMPLE 132

6-(4-Fluorophenyl)-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine-3-carbonitrile

Following a similar procedure to that described in example 65, but using 3-bromo-6-(4-fluorophenyl)-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine (obtained in example 76) instead of 3-bromo-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine, the title compound was obtained (yield: 38%).

EXAMPLE 133

2-[2-[[1-(*tert*-Butoxycarbonyl)piperidin-4-yl]amino]ethyl]-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine

Following a similar procedure to that described in example 111, but using 2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl methanesulfonate (obtained in example 116 section a) instead of 3-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]propyl methanesulfonate and [N-(*tert*-butoxycarbonyl)piperidin-4-yl]amine instead of N-(*tert*-butoxycarbonyl)-N-(4-piperidyl)amine, the desired compound was obtained (yield: 40%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.20 (m, 2 H), 1.45 (s, 9 H), 1.55 (broad s NH + H₂O), 1.83 (m, 2 H), 2.65 (m, 1 H), 2.77 (m, 2 H), 3.31 (t, 2 H), 4.02 (m, 2 H), 4.53 (t, 2 H), 6.82 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H), 6.90 (t, J = 8.8 Hz, 2 H), 7.00 (t, J = 8.7 Hz, 2 H), 7.13 (m, 2 H), 7.28 (m, 2 H), 7.84 (s, 1 H), 8.32 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H).

EXAMPLE 134

4,6-Bis(4-fluorophenyl)-2-[2-[(4-piperidyl)amino]ethyl]-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine

Following a similar procedure to that described in example 36, but using 2-[2-[[1-(*tert*-butoxycarbonyl)piperidin-4-yl]amino]ethyl]-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine (obtained in example 133) instead of 2-[1-(*tert*-butoxycarbonyl)piperidin-4-yl]-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine, the title compound was obtained (yield: 97%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.30 (m, 2 H), 1.70 (broad s, 2 NH + H₂O), 1.85 (m, 2 H), 2.61 (m, 3 H), 3.09 (m, 2 H), 3.31 (t, J = 5.7 Hz, 2 H), 4.53 (t, J = 5.7 Hz, 2 H), 6.82 (dd, J_o = 1.6 Hz, J_m = 4.4 Hz, 2 H), 6.90 (t, J = 8.8 Hz, 2 H), 7.00 (t, J = 8.7 Hz, 2 H), 7.14 (m, 2 H), 7.29 (m, 2 H), 7.85 (s, 1 H), 8.32 (dd, J_o = 1.6 Hz, J_m = 4.4 Hz, 2 H).

EXAMPLE 135

N-(2-Methoxyethyl)-[2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl]amine

Following a similar procedure to that described in example 111, but using 2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl methanesulfonate (obtained in example 116 section a) instead of 3-[4,6-bis(4-fluorophenyl)-5-(4-

pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]propyl methanesulfonate and 2-methoxyethylamine instead of *N*-(*tert*-butoxycarbonyl)-*N*-(4-piperidyl)amine, the desired compound was obtained (yield: 63%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.57 (broad s, NH + H₂O), 2.81 (t, J = 5.1 Hz, 2 H), 3.30 (t, J = 5.8 Hz, 2 H), 3.31 (s, 3 H), 3.45 (t, J = 5.1 Hz, 2 H), 4.55 (t, J = 5.7 Hz, 2 H), 6.82 (dd, J_o = 1.8 Hz, J_m = 4.5 Hz, 2 H), 6.89 (t, J = 8.7 Hz, 2 H), 7.00 (t, J = 8.7 Hz, 2 H), 7.14 (m, 2 H), 7.29 (m, 2 H), 7.85 (s, 1 H), 8.32 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H).

EXAMPLE 136

10 1-[4-[2-[4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl]piperazin-1-yl]ethanone

Following a similar procedure to that described in example 111, but using 2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl methanesulfonate (obtained in example 116 section a) instead of 3-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]propyl methanesulfonate and 1-(piperazin-1-yl)ethanone instead of *N*-(*tert*-butoxycarbonyl)-*N*-(4-piperidyl)amine, the desired compound was obtained (yield: 50%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 2.07 (s, 3 H), 2.50 (m, 4 H), 3.07 (t, J = 6.3 Hz, 2 H), 3.41 (t, J = 4.9 Hz, 2 H), 3.57 (t, J = 4.9 Hz, 2 H), 4.55 (t, J = 6.3 Hz, 2 H), 6.82 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H), 6.90 (t, J = 8.7 Hz, 2 H), 7.00 (t, J = 8.6 Hz, 2 H), 7.14 (m, 2 H), 7.29 (m, 2 H), 7.82 (s, 1 H), 8.32 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H).

EXAMPLE 137

3-[4,6-Diphenyl-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]propan-1-ol

25

EXAMPLE 138

3-[4,6-Diphenyl-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-1-yl]propan-1-ol

Following a similar procedure to that described in examples 6 and 7, but using 4,6-diphenyl-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine (obtained in example 2) instead of 4,6-bis(4-fluorophenyl)-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine and 3-iodopropanol instead of iodoethane, the title compounds were obtained.

Example 137: yield: 44%; ¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.56 (s, OH + H₂O), 2.25 (m, 2 H), 3.70 (m, 2 H), 4.62 (t, J = 6.3 Hz, 2 H), 6.83 (dd, J_o = 1.6 Hz,

$J_m = 4.6$ Hz, 2 H), 7.17 – 7.34 (complex-signal, 10 H), 7.84 (s, 1 H), 8.26 (dd, $J_o = 1.5$ Hz, $J_m = 4.5$ Hz, 2 H).

Example 138: yield: 27%; ^1H NMR (300 MHz, CDCl_3) δ (TMS): 1.58 (s, OH + H_2O), 2.15 (m, 2 H), 3.59 (m, 2 H), 4.79 (t, $J = 6.0$ Hz, 2 H), 6.83 (dd, $J_o = 1.6$ Hz, $J_m = 4.4$ Hz, 2 H), 7.17 (m, 2 H), 7.21 – 7.33 (complex signal, 8 H), 7.91 (s, 1 H), 8.27 (dd, $J_o = 1.5$ Hz, $J_m = 4.5$ Hz, 2 H).

EXAMPLE 139

2-Ethyl-4,6-diphenyl-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine

EXAMPLE 140

10 1-Ethyl-4,6-diphenyl-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine

Following a similar procedure to that described in examples 6 and 7, but using 4,6-diphenyl-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine (obtained in example 2) instead of 4,6-bis(4-fluorophenyl)-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine, the title compounds were obtained.

15 Example 139: yield: 12%; ^1H NMR (300 MHz, CDCl_3) δ (TMS): 1.67 (t, $J = 7.3$ Hz, 3 H), 4.50 (c, $J = 7.3$ Hz, 2 H), 6.83 (dd, $J_o = 1.5$ Hz, $J_m = 4.5$ Hz, 2 H), 7.15 – 7.34 (complex signal, 10 H), 7.80 (s, 1H), 8.25 (dd, $J_o = 1.5$ Hz, $J_m = 4.5$ Hz, 2 H).

Example 140: yield: 21%; ^1H NMR (300 MHz, CDCl_3) δ (TMS): 1.61 (t, $J = 7.2$ Hz, 3 H), 4.68 (c, $J = 7.2$ Hz, 2 H), 6.82 (dd, $J_o = 1.5$ Hz, $J_m = 4.5$ Hz, 2 H), 7.17 (m, 2 H), 7.23 – 7.32 (complex signal, 8 H), 7.88 (s, 1H), 8.26 (dd, $J_o = 1.5$ Hz, $J_m = 4.5$ Hz, 2 H).

EXAMPLE 141

4,6-Diphenyl-2-(2-phthalimidoethyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine

Following a similar procedure to that described in examples 6 and 7, but using 4,6-diphenyl-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine (obtained in example 2) instead of 4,6-bis(4-fluorophenyl)-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine and *N*-(2-bromoethyl)phthalimide instead of iodoethane, the title compound was obtained (yield: 31%).

25 ^1H NMR (300 MHz, CDCl_3) δ (TMS): 4.32 (t, $J = 6.1$ Hz, 2 H), 4.75 (t, $J = 6.1$ Hz, 2 H), 6.82 (dd, $J_o = 1.5$ Hz, $J_m = 4.5$ Hz, 2 H), 7.11 (m, 2 H), 7.18 – 7.32 (complex signal, 7 H), 7.73 (m, 2 H), 7.80 (s, 1 H), 7.83 (m, 2 H), 8.25 (dd, $J_o = 1.5$ Hz, $J_m = 4.5$ Hz, 2 H).

EXAMPLE 142

2-(2-Aminoethyl)-4,6-diphenyl-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine

Following a similar procedure to that described in example 90, but using 2-(2-phthalimidoethyl)-4,6-diphenyl-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine (obtained in example 141) instead of 4,6-bis(4-fluorophenyl)-2-(2-phthalimidoethyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine, the title compound was obtained (yield: 51%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.50 (s, NH₂ + H₂O), 3.37 (t, J = 5.5 Hz, 2 H), 4.48 (t, J = 5.4 Hz, 2 H), 6.83 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H), 7.17 – 7.34 (complex signal, 10 H), 7.86 (s, 1 H), 8.25 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H).

EXAMPLE 143

10 **2-Allyl-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine**

EXAMPLE 144**1-Allyl-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine**

Following a similar procedure to that described in examples 6 and 7, but using allyl bromide instead of iodoethane, the title compounds were obtained.

15 Example 143: yield: 33%; ¹H NMR (300 MHz, CDCl₃) δ (TMS): 5.08 (d, J = 6.3 Hz, 2 H), 5.40 (m, 2 H), 6.16 (m, 1 H), 6.82 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H), 6.89 (t, J = 8.7 Hz, 2 H), 7.00 (t, J = 8.6 Hz, 2 H), 7.13 (m, 2 H), 7.30 (m, 2 H), 7.78 (s, 1 H), 8.34 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H).

Example 144: yield: 10%; ¹H NMR (300 MHz, CDCl₃) δ (TMS): 5.22 – 5.34 (complex signal, 4 H), 6.14 (m, 1 H), 6.81 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H), 6.93 (t, J = 8.7 Hz, 2 H), 7.01 (t, J = 8.6 Hz, 2 H), 7.14 (m, 2 H), 7.29 (m, 2 H), 7.87 (s, 1 H), 8.32 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H).

EXAMPLE 145

25 **1-[2-[4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl]piperidin-4-one**

Following a similar procedure to that described in example 111, but using 2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl methanesulfonate (obtained in example 116 section a) instead of 3-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]propyl methanesulfonate and 4-piperidone monohydrate hydrochloride instead of *N*-(*tert*-butoxycarbonyl)-*N*-(4-piperidyl)amine and adding triethylamine (3 equivalents), the desired product was obtained (yield: 18%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 2.41 (t, J = 6.0 Hz, 4 H), 2.84 (t, J = 6.0 Hz, 4 H), 3.20 (t, J = 6.4 Hz, 2 H), 4.58 (t, J = 6.4 Hz, 2 H), 6.83 (d, J = 6.0 Hz, 2 H), 6.90 (t, J = 8.7 Hz, 2 H), 7.00 (t, J = 8.6 Hz, 2 H), 7.14 (m, 2 H), 7.30 (m, 2 H), 7.85 (s, 1 H), 8.32 (d, J = 6.0 Hz, 2 H).

5

EXAMPLE 146**3-Aminomethyl-6-(4-fluorophenyl)-5-(4-pyridyl)-1H-pyrazolo[3,4-b]pyridine**

Following a similar procedure to that described in example 68, but using 6-(4-fluorophenyl)-5-(4-pyridyl)-1H-pyrazolo[3,4-b]pyridine-3-carbonitrile (obtained in example 132) instead of 4,6-bis(4-fluorophenyl)-5-(4-pyridyl)-1H-pyrazolo[3,4-

10

b]pyridine-3-carbonitrile, the title compound was obtained (yield: 22%).
¹H NMR (300 MHz, CDCl₃ + CD₃OD) δ (TMS): 4.24 (s, 2 H), 4.25 (broad s, NH + NH₂ + CD₃OD), 6.95 (m, 2 H), 7.16 (m, 2 H), 7.29 (m, 2 H), 8.25 (broad s, 1 H), 8.38 (broad s, 2 H).

EXAMPLE 147**15 3-Amino-6-(4-fluorophenyl)-4-methyl-5-(4-pyridyl)-1H-pyrazolo[3,4-b]pyridine**

Following a similar procedure to that described in reference example 15 section c, but using 6-chloro-2-(4-fluorophenyl)-4-methyl-3,4'-bipyridine-5-carbonitrile (obtained in reference example 18) instead of 3-(1-benzylpiperidin-4-yl)-3-oxopropiononitrile, the title compound was obtained (yield: 27%).

20 ¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.56 (broad s, NH + NH₂ + H₂O), 2.51 (s, 3 H), 6.80 – 7.20 (complex signal, 4 H), 7.22 (m, 2 H), 8.55 (d, J = 8.0 Hz, 2 H).

EXAMPLE 148**3-[N-[2-[4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridin-2-yl]ethyl]amino]propan-1-ol**

25 Following a similar procedure to that described in example 111, but using 2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridin-2-yl]ethyl methanesulfonate (obtained in example 116 section a) instead of 3-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridin-2-yl]propyl methanesulfonate and 3-amino-1-propanol instead of N-(tert-butoxycarbonyl)-N-(4-piperidyl)amine, the desired
30 compound was obtained (yield: 57%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.50 – 1.80 (complex signal, 2 H + NH + OH + H₂O), 2.91 (t, J = 5.7 Hz, 2 H), 3.31 (t, J = 5.4 Hz, 2 H), 3.77 (t, J = 5.4 Hz, 2 H), 4.54 (t, J = 5.4 Hz, 2 H), 6.82 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H), 6.90 (t, J = 8.7

Hz, 2 H), 7.00 (t, $J = 8.7$ Hz, 2 H), 7.15 (m, 2 H), 7.29 (m, 2 H), 7.82 (s, 1 H), 8.31 (dd, $J_o = 1.8$ Hz, $J_m = 4.5$ Hz, 2 H).

EXAMPLE 149

N-Ethyl-[2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl]amine

5 Following a similar procedure to that described in example 111, but using 2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl methanesulfonate (obtained in example 116 of section a) instead of 3-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]propyl methanesulfonate and ethylamine instead
10 of *N*-(*tert*-butoxycarbonyl)-*N*-(4-piperidyl)amine, the desired compound was obtained (yield: 58%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.09 (t, $J = 7.0$ Hz, 3 H), 1.71 (broad s, NH + H₂O), 2.70 (c, $J = 7.1$ Hz, 2 H), 3.29 (t, $J = 5.7$ Hz, 2 H), 4.56 (t, $J = 5.7$ Hz, 2 H), 6.82 (dd, $J_o = 1.5$ Hz, $J_m = 4.5$ Hz, 2 H), 6.90 (t, $J = 8.7$ Hz, 2 H), 7.00 (t, $J = 8.7$
15 Hz, 2 H), 7.14 (m, 2 H), 7.29 (m, 2 H), 7.84 (s, 1 H), 8.32 (dd, $J_o = 1.5$ Hz, $J_m = 4.5$ Hz, 2 H).

EXAMPLE 150

2-[N-[2-[4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl]amino]ethanol

20 Following a similar procedure to that described in example 111, but using 2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl methanesulfonate (obtained in example 116 section a) instead of 3-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]propyl methanesulfonate and 2-aminoethanol instead of *N*-(*tert*-butoxycarbonyl)-*N*-(4-piperidyl)amine, the desired compound
25 was obtained (yield: 54%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.70 (broad s, NH + OH + H₂O), 2.82 (t, $J = 5.3$ Hz, 2 H), 3.32 (t, $J = 5.6$ Hz, 2 H), 3.62 (t, $J = 5.3$ Hz, 2 H), 4.55 (t, $J = 5.6$ Hz, 2 H), 6.83 (dd, $J_o = 1.5$ Hz, $J_m = 4.5$ Hz, 2 H), 6.90 (t, $J = 8.7$ Hz, 2 H), 7.00 (t, $J = 8.6$ Hz, 2 H), 7.14 (m, 2 H), 7.29 (m, 2 H), 7.83 (s, 1 H), 8.31 (dd, $J_o = 1.8$ Hz, $J_m =$
30 4.5 Hz, 2 H).

EXAMPLE 151

N-[(2-Pyridyl)methyl]-[2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl]amine

Following a similar procedure to that described in example 111, but using 2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl methanesulfonate (obtained in example 116 section a) instead of 3-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]propyl methanesulfonate and (2-pyridyl)methylamine instead of *N*-(*tert*-butoxycarbonyl)-*N*-(4-piperidyl)amine, the
5 desired compound was obtained (yield: 52%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.62 (broad s, NH + H₂O), 3.33 (t, J = 5.7 Hz, 2 H), 3.92 (s, 2 H), 4.58 (t, J = 5.7 Hz, 2 H), 6.83 (dd, J_o = 1.8 Hz, J_m = 4.5 Hz, 2 H), 6.90 (t, J = 8.7 Hz, 2 H), 6.99 (t, J = 8.6 Hz, 2 H), 7.14 (m, 4 H), 7.29 (m, 2 H),
10 7.60 (m, 1 H), 7.86 (s, 1 H), 8.31 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H), 8.51 (m, 1 H).

EXAMPLE 152

N-[(2-Thienyl)methyl]-[2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl]amine

Following a similar procedure to that described in example 111, but using 2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl methanesulfonate (obtained in example 116 section a) instead of 3-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]propyl and (2-thienyl)methylamine instead of *N*-(*tert*-butoxycarbonyl)-*N*-(4-piperidyl)amine, the desired compound was obtained
15 (yield: 25%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.55 (broad s, NH + H₂O) 3.32 (t, J = 5.6 Hz, 2 H), 4.00 (s, 2 H), 4.54 (t, J = 5.6 Hz, 2 H), 6.83 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H), 6.90 (t, J = 8.7 Hz, 2 H), 6.91 (m, 1 H), 7.00 (t, J = 8.6 Hz, 2 H), 7.16 (m, 3 H), 7.29 (m, 3 H), 7.86 (s, 1 H), 8.32 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H).

EXAMPLE 153

1-[2-[4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl]piperidine-4-carboxamide

Following a similar procedure to that described in example 111, but using 2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl methanesulfonate (obtained in example 116 section a) instead of 3-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]propyl methanesulfonate and piperidine-4-carboxamide instead of *N*-(*tert*-butoxycarbonyl)-*N*-(4-piperidyl)amine, the desired
30 compound was obtained (yield: 75%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.61 – 1.76 (complex signal, 5 H), 1.84 (m, 2 H), 2.91 (m, 2 H), 3.01 (t, J = 6.3 Hz, 2 H), 4.52 (t, J = 6.3 Hz, 2 H), 5.27 (broad s, NH), 5.41 (broad s, NH), 6.83 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H), 6.89 (t, J = 8.7 Hz, 2 H), 7.02 (t, J = 8.6 Hz, 2 H), 7.14 (m, 2 H), 7.30 (m, 2 H), 7.83 (s, 1 H), 8.32 (dd, J_o = 1.4 Hz, J_m = 4.6 Hz, 2 H).

EXAMPLE 154

4,6-Bis(4-fluorophenyl)-2-[2-(pyrrolidin-1-yl)ethyl]-5-(4-pyridyl)pyrazolo[3,4-b]pyridine

Following a similar procedure to that described in example 111, but using 2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridin-2-yl]ethyl methanesulfonate (obtained in example 116 section a) instead of 3-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridin-2-yl]propyl methanesulfonate and pyrrolidine instead of *N*-(*tert*-butoxycarbonyl)-*N*-(4-piperidyl)amine, the desired compound was obtained (yield: 75%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.77 (m, 4 H), 2.55 (m, 4 H), 3.17 (t, J = 6.6 Hz, 2 H), 4.58 (t, J = 6.6 Hz, 2 H), 6.82 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H), 6.89 (t, J = 8.7 Hz, 2 H), 7.00 (t, J = 8.7 Hz, 2 H), 7.14 (m, 2 H), 7.29 (m, 2 H), 7.84 (s, 1 H), 8.31 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H).

EXAMPLE 155

(3*R*)-1-[2-[4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridin-2-yl]ethyl]pyrrolidin-3-ol

Following a similar procedure to that described in example 111, but using 2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridin-2-yl]ethyl methanesulfonate (obtained in example 116 section a) instead of 3-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridin-2-yl]propyl methanesulfonate and (3*R*)-3-pyrrolidinol instead of *N*-(*tert*-butoxycarbonyl)-*N*-(4-piperidyl)amine, the desired compound was obtained (yield: 52%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.69 (broad s, 1H + OH + H₂O), 2.14 (m, 1 H), 2.40 (m, 1 H), 2.58 (m, 1 H), 2.69 (m, 1 H), 2.94 (m, 1 H), 3.20 (t, J = 6.4 Hz, 2 H), 4.32 (m, 1 H), 4.57 (t, J = 6.4 Hz, 2 H), 6.83 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H), 6.89 (t, J = 8.7 Hz, 2 H), 7.00 (t, J = 8.6 Hz, 2 H), 7.14 (m, 2 H), 7.29 (m, 2 H), 7.82 (s, 1 H), 8.32 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H).

EXAMPLE 156

2-[N-[2-[4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridin-2-yl]ethyl]-N-methylamino]ethanol

Following a similar procedure to that described in example 111, but using 2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridin-2-yl]ethyl methanesulfonate (obtained in example 116 section a) instead of 3-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridin-2-yl]propyl methanesulfonate and 2-(methylamino)ethanol instead of *N*-(*tert*-butoxycarbonyl)-*N*-(4-piperidyl)amine, the desired compound was obtained (yield: 71%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.56 (broad s, OH + H₂O), 2.34 (s, 3 H), 2.60 (t, J = 5.2 Hz, 2 H), 3.14 (t, J = 6.0 Hz, 2 H), 3.53 (t, J = 5.2 Hz, 2 H), 4.53 (t, J = 6.0 Hz, 2 H), 6.83 (dd, J_o = 1.8 Hz, J_m = 4.5 Hz, 2 H), 6.89 (t, J = 8.7 Hz, 2 H), 7.00 (t, J = 8.6 Hz, 2 H), 7.14 (m, 2 H), 7.29 (m, 2 H), 7.81 (s, 1 H), 8.31 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H).

EXAMPLE 157

4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)-2-[2-(1,2,3,4-tetrahydroisoquinolin-2-yl)ethyl]pyrazolo[3,4-b]pyridine

Following a similar procedure to that described in example 111, but using 2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridin-2-yl]ethyl methanesulfonate (obtained in example 116 section a) instead of 3-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridin-2-yl]propyl methanesulfonate and 1,2,3,4-tetrahydroisoquinoline instead of *N*-(*tert*-butoxycarbonyl)-*N*-(4-piperidyl)amine, the desired compound was obtained (yield: 57%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 2.78 – 2.85 (complex signal, 4 H), 3.20 (t, J = 6.3 Hz, 2 H), 3.72 (s, 2 H), 4.63 (t, J = 6.3 Hz, 2 H), 6.81 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H), 6.86 – 7.17 (complex signal, 10 H), 7.30 (m, 2 H), 7.86 (s, 1 H), 8.31 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H).

EXAMPLE 158

4,6-Bis(4-fluorophenyl)-2-[2-(4-phenylpiperazin-1-yl)ethyl]-5-(4-pyridyl)pyrazolo[3,4-b]pyridine

Following a similar procedure to that described in example 111, but using 2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridin-2-yl]ethyl methanesulfonate (obtained in example 116 section a) instead of 3-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridin-2-yl]propyl methanesulfonate and 1-

phenylpiperazine instead of *N*-(*tert*-butoxycarbonyl)-*N*-(4-piperidyl)amine, the desired compound was obtained (yield: 71%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 2.69 (m, 4 H), 3.10 (t, J = 6.4 Hz, 2 H), 3.16 (m, 4 H), 4.59 (t, J = 6.4 Hz, 2 H), 6.82 (dd, J_o = 1.8 Hz, J_m = 4.5 Hz, 2 H), 6.86 – 7.02 (complex signal, 9 H), 7.14 (m, 2 H), 7.29 (m, 2 H), 7.86 (s, 1 H), 8.32 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H).

EXAMPLE 159

4,6-Bis(4-fluorophenyl)-2-[2-[4-(1-piperidyl)piperidin-1-yl]ethyl]-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine

10 Following a similar procedure to that described in example 111, but using 2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl methanesulfonate (obtained in example 116 section a) instead of 3-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]propyl methanesulfonate and 4-piperidinopiperidine instead of *N*-(*tert*-butoxycarbonyl)-*N*-(4-piperidyl)amine, the
15 desired compound was obtained (yield: 51%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.40 – 1.60 (complex signal, 8 H), 1.78 (m, 2 H), 2.04 – 2.22 (complex signal, 3 H), 2.48 (m, 4 H), 2.96 (m, 4 H), 4.52 (t, J = 6.4 Hz, 2 H), 6.82 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H), 6.89 (t, J = 8.7 Hz, 2 H), 7.00 (t, J = 8.6 Hz, 2 H), 7.13 (m, 2 H), 7.29 (m, 2 H), 7.85 (s, 1 H), 8.31 (dd, J_o = 1.5 Hz,
20 J_m = 4.5 Hz, 2 H).

EXAMPLE 160

3-[*N*-2-[4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl]-*N*-methylamino]propiononitrile

Following a similar procedure to that described in example 111, but using 2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl methanesulfonate
25 (obtained in example 116 section a) instead of 3-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]propyl methanesulfonate and 3-(methylamino)propiononitrile instead of *N*-(*tert*-butoxycarbonyl)-*N*-(4-piperidyl)amine, the desired compound was obtained (yield: 50%).

30 ¹H NMR (300 MHz, CDCl₃) δ (TMS): 2.35 (complex signal, 5 H), 2.71 (t, J = 6.4 Hz, 4 H), 3.13 (t, J = 5.9 Hz, 2 H), 4.50 (t, J = 5.9 Hz, 2 H), 6.83 (dd, J_o = 1.6 Hz, J_m = 4.4 Hz, 2 H), 6.90 (t, J = 8.7 Hz, 2 H), 6.99 (t, J = 8.7 Hz, 2 H), 7.17 (m, 2 H), 7.31 (m, 2 H), 7.87 (s, 1 H), 8.32 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H).

EXAMPLE 161***N*-Methyl-2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethylamine**

Following a similar procedure to that described in example 111, but using 2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl methanesulfonate (obtained in example 116 section a) instead of 3-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]propyl methanesulfonate and methylamine instead of *N*-(*tert*-butoxycarbonyl)-*N*-(4-piperidyl)amine, the desired compound was obtained (yield: 51%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.61 (broad s, NH + H₂O), 3.25 (t, J = 5.6 Hz, 2 H), 4.57 (t, J = 5.6 Hz, 2 H), 6.82 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H), 6.90 (t, J = 8.7 Hz, 2 H), 7.00 (t, J = 8.6 Hz, 2 H), 7.14 (m, 2 H), 7.29 (m, 2 H), 7.84 (s, 1 H), 8.31 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H).

EXAMPLE 162**2-[2-[4-(*tert*-Butoxycarbonyl)piperazin-1-yl]ethyl]-4,6-(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine**

Following a similar procedure to that described in example 111, but using 2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl methanesulfonate (obtained in example 116 section a) instead of 3-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]propyl methanesulfonate and 1-(*tert*-butoxycarbonyl)piperazine instead of *N*-(*tert*-butoxycarbonyl)-*N*-(4-piperidyl)amine, the desired compound was obtained (yield: 63%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.45 (s, 9 H), 2.46 (m, 4 H), 3.05 (t, J = 6.4 Hz, 2 H), 3.40 (m, 4 H), 4.54 (t, J = 6.4 Hz, 2 H), 6.82 (dd, J_o = 1.6 Hz, J_m = 4.4 Hz, 2 H), 6.89 (t, J = 8.7 Hz, 2 H), 7.00 (t, J = 8.6 Hz, 2 H), 7.15 (m, 2 H), 7.30 (m, 2 H), 7.86 (s, 1 H), 8.33 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H).

EXAMPLE 163**4,6-Bis(4-fluorophenyl)-2-[2-(piperazin-1-yl)ethyl]-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine**

Following a similar procedure to that described in example 36, but using 2-[2-[4-(*tert*-butoxycarbonyl)piperazin-1-yl]ethyl]-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine (obtained in example 162) instead of 2-[1-(*tert*-butoxycarbonyl)piperidin-4-yl]-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-

b]pyridine, the title compound was obtained (yield: 62%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.56 (broad s, NH + H₂O), 2.45 (m, 4 H), 2.85 (m, 4 H), 3.00 (t, J = 6.4 Hz, 2 H), 4.55 (t, J = 6.4 Hz, 2 H), 6.82 (dd, J_o = 1.6 Hz, J_m = 4.4 Hz, 2 H), 6.89 (t, J = 8.7 Hz, 2 H), 7.03 (t, J = 8.7 Hz, 2 H), 7.14 (m, 2 H),
5 7.30 (m, 2 H), 7.86 (s, 1 H), 8.32 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H).

EXAMPLE 164

4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)-2-vinylpyrazolo[3,4-b]pyridine

A solution of 2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridin-2-yl]ethyl methanesulfonate (0.15 g, 0.3 mmol, obtained in example 116 section a) and KOH (0.02 g, 0.3 mmol) in toluene (4 mL) was heated to 100 °C reflux overnight. Water and EtOAc were added and the phases were separated. The aqueous phase was
10 saturated with NaCl (solid) and extracted with EtOAc. The combined organic phases were dried over Na₂SO₄ and concentrated to dryness. The crude product obtained was purified by chromatography on silica gel using hexane-EtOAc
15 mixtures of increasing polarity as eluent, to afford 110 mg of the title compound (yield: 90%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 5.27 (dd, J_{gem} = 1.8 Hz, J_{vec} = 8.7 Hz, 1 H), 6.22 (dd, J_{gem} = 1.4 Hz, J_{vec} = 15.4 Hz, 1 H), 6.83 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H), 6.91 (t, J = 8.7 Hz, 2 H), 7.01 (t, J = 8.6 Hz, 2 H), 7.15 (m, 2 H), 7.17 – 7.34
20 (complex signal, 3 H), 7.90 (s, 1 H), 8.33 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H).

EXAMPLE 165

2-[N-[2-[4,6-Bis-(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridin-2-yl]ethyl]-N-(2-hydroxyethyl)amino]ethanol

Following a similar procedure to that described in example 111, but using 2-[4,6-bis-(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridin-2-yl]ethyl methanesulfonate
25 (obtained in example 116 section a) instead of 3-[4,6-bis-(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridin-2-yl]propyl methanesulfonate and 2-(2-hydroxyethylamino)ethanol instead of N-(tert-butoxycarbonyl)-N-(4-piperidyl)amine, the desired compound was obtained (yield: 51%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.50 (broad s, 2 OH + H₂O), 2.73 (t, J = 5.0 Hz, 4H), 3.21 (t, J = 5.6 Hz, 2 H), 3.52 (t, J = 5.0 Hz, 4 H), 4.54 (t, J = 5.4 Hz, 2 H), 6.82 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H), 6.89 (t, J = 8.7 Hz, 2 H), 6.99 (t, J = 8.7

Hz, 2 H), 7.13 (m, 2 H), 7.29 (m, 2 H), 7.89 (s, 1 H), 8.32 (dd, $J_o = 1.8$ Hz, $J_m = 4.5$ Hz, 2 H).

EXAMPLE 166

***N*-Cyclopropyl-2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethylamine**

Following a similar procedure to that described in example 111, but using 2-[4,6-bis-(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl methanesulfonate (obtained in example 116 section a) instead of 3-[4,6-bis-(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]propyl methanesulfonate and cyclopropylamine instead of *N*-(*tert*-butoxycarbonyl)-*N*-(4-piperidyl)amine, the desired compound was obtained (yield: 47%).

^1H NMR (300 MHz, CDCl_3) δ (TMS): 0.30 (m, 2 H), 0.45 (m, 2 H), 1.60 (broad s, NH + H_2O), 2.20 (m, 1 H), 3.36 (t, $J = 5.7$ Hz, 2 H), 4.55 (t, $J = 5.7$ Hz, 2 H), 6.82 (dd, $J_o = 1.5$ Hz, $J_m = 4.5$ Hz, 2 H), 6.90 (t, $J = 8.7$ Hz, 2 H), 7.00 (t, $J = 8.6$ Hz, 2 H), 7.14 (m, 2 H), 7.30 (m, 2 H), 7.82 (s, 1 H), 8.32 (dd, $J_o = 1.5$ Hz, $J_m = 4.5$ Hz, 2 H).

EXAMPLE 167

***N*-[2-[4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl]acetamide**

Following a similar procedure to that described in reference example 1 section a, but using acetyl chloride instead of 4-fluorobenzoyl chloride and 2-(2-aminoethyl)-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine (obtained in example 90) instead of *N,O*-dimethylhydroxylamine, the desired compound was obtained (yield: 48%).

^1H NMR (300 MHz, CDCl_3) δ (TMS): 1.98 (s, 3 H), 3.95 (m, 2 H), 4.58 (t, $J = 5.4$ Hz, 2 H), 6.47 (m, NH), 6.83 (d, $J = 9.0$ Hz, 2 H), 6.91 (t, $J = 8.7$ Hz, 2 H), 7.01 (t, $J = 8.6$ Hz, 2 H), 7.14 (m, 2 H), 7.28 (m, 2 H), 7.82 (s, 1 H), 8.33 (d, $J = 9.0$ Hz, 2 H).

EXAMPLE 168

***N*-[2-[4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl]-*N'*-isopropylurea**

To a solution of 2-(2-aminoethyl)-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine (0.06 g, 0.15 mmol, obtained in example 90) in DMF (1 mL), isopropylisocyanate (0.02 g, 0.18 mmol) was added under argon atmosphere. This

was stirred at room temperature for 2 days. The solvent was concentrated again and diethyl ether was added to the residue obtained. The solvent was concentrated, to afford 38 mg of title compound in solid form (yield: 50%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.05 (d, J = 6.3 Hz, 6 H), 3.84 (m, 3 H), 4.22 (m, NH), 4.58 (t, J = 5.4 Hz, 2 H), 5.30 (m, NH), 6.82 (dd, J_o = 1.4 Hz, J_m = 4.6 Hz, 2 H), 6.91 (t, J = 8.7 Hz, 2 H), 7.00 (t, J = 8.6 Hz, 2 H), 7.13 (m, 2 H), 7.28 (m, 2 H), 7.84 (s, 1 H), 8.32 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H).

EXAMPLE 169

N-[2-[4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl]methanesulfonamide

To a solution of 2-(2-aminoethyl)-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine (0.06 g, 0.15 mmol, obtained in example 90) and DMAP (0.001 g, 0.0058 mmol) in pyridine (0.6 mL), methanesulfonyl chloride (0.017 mL, 0.22 mmol) was added under argon atmosphere and cooled with an ice bath (0.017 mL, 0.22 mmol). This was stirred overnight at room temperature. The solvent was concentrated. The residue was dissolved in CHCl₃ and saturated NaHCO₃ was added. The phases were separated. The organic phase was dried over Na₂SO₄ and concentrated. The crude product obtained was purified by chromatography on silica gel using EtOAc as solvent, and 70 mg of the title compound was obtained (yield: 95%).

¹H NMR (300 MHz, CDCl₃) δ (TMS): 2.99 (s, 3 H), 3.86 (m, 2 H), 4.64 (t, J = 5.3 Hz, 2 H), 5.33 (m, NH), 6.83 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H), 6.91 (t, J = 8.7 Hz, 2 H), 7.01 (t, J = 8.7 Hz, 2 H), 7.14 (m, 2 H), 7.29 (m, 2 H), 7.88 (s, 1 H), 8.33 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H).

EXAMPLES 170-178

Following a similar procedure to that described in example 72, but starting from the appropriate compounds in each case, the compounds in the following table were obtained:

Example	Compound name	Starting compounds	LC-MS		
			Method	t _R (min)	m/z [M+H] ⁺
170	6-(4-Fluorophenyl)-4-(4-piperidyl)-5-(4-pyridyl)-1H-	Reference example 1, 3-amino-2H-pyrazole and	1	3.34	374.1

	pyrazolo[3,4- <i>b</i>]pyridine ..	1-(<i>tert</i> -butoxycarbonyl)- piperidine-4- carbaldehyde			
171	6-(4-Fluorophenyl)-4-(2-furyl)-5-(4-pyridyl)-1 <i>H</i> -pyrazolo[3,4- <i>b</i>]pyridine	Reference example 1, 3-amino-2 <i>H</i> -pyrazole and furan-2-carbaldehyde	1	5.72	357.1
172	6-(4-Fluorophenyl)-4-(1 <i>H</i> -imidazol-4-yl)-5-(4-pyridyl)-1 <i>H</i> -pyrazolo[3,4- <i>b</i>]pyridine	Reference example 1, 3-amino-2 <i>H</i> -pyrazole and 2 <i>H</i> -pyrazole-3-carbaldehyde	1	3.25	357.1
173	4-(5-Bromothiophen-2-yl)-6-(4-fluorophenyl)-5-(4-pyridyl)-1 <i>H</i> -pyrazolo[3,4- <i>b</i>]pyridine	Reference example 1, 3-amino-2 <i>H</i> -pyrazole and 5-bromothiophene-2-carbaldehyde	1	7.46	450.9 452.9
174	4,6-Bis(4-fluorophenyl)-5-(2-methylsulfanylpirimidin-4-yl)-1 <i>H</i> -pyrazolo[3,4- <i>b</i>]pyridine	Reference example 19, 3-amino-2 <i>H</i> -pyrazole and 4-fluorobenzaldehyde	1	9.11	432.2
175	5-(2-Chloropyridin-4-yl)-4,6-bis(4-fluorophenyl)-1 <i>H</i> -pyrazolo[3,4- <i>b</i>]pyridine	Reference example 23, 3-amino-2 <i>H</i> -pyrazole and 4-fluorobenzaldehyde	1	8.99	419.0 421.0
176	6-(4-Fluorophenyl)-4-(2-phenylethyl)-5-(4-pyridyl)-1 <i>H</i> -pyrazolo[3,4- <i>b</i>]pyridine	Reference example 1, 3-amino-2 <i>H</i> -pyrazole and 3-phenylpropionaldehyde	1	6.52	395.0
177	4-(6-Chloropyridin-3-yl)-6-(4-fluorophenyl)-5-(4-pyridyl)-1 <i>H</i> -pyrazolo[3,4- <i>b</i>]pyridine	Reference example 1, 6-chloropyridine-3-carbaldehyde and amino-2 <i>H</i> -pyrazole	1	5.50	402.0 404.0
178	4-(3,4-Dichlorophenyl)-1-ethyl-6-(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4- <i>b</i>]pyridine	Reference example 1, 3,4-dichlorobenzaldehyde and 3-amino-2-ethylpyrazole	1	10.56	462.9 464.9

EXAMPLE 179

6-(4-Fluorophenyl)-4-(1-methylpiperidin-4-yl)-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine

To a suspension of 6-(4-fluorophenyl)-4-(4-piperidyl)-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine (0.24 g, 0.6 mmol, obtained in example 170) in formic acid (0.64 mL), 35-40 % aqueous formaldehyde (0.96 mL) was added. It was heated to 70-80 °C for 24 h. It was allowed to cool and 1 N NaOH was added. It was extracted with CHCl_3 and the combined organic phases were dried over Na_2SO_4 and concentrated to dryness. The crude product obtained was purified by chromatography on silica gel using CHCl_3 -MeOH- NH_3 mixtures of increasing polarity as eluent, to afford 47 mg of the desired compound (yield: 19%).

LC-MS (method 1): t_R = 3.28 min; m/z = 388.1 $[\text{M}+\text{H}]^+$.

10

EXAMPLE 180**3-Amino-6-(4-fluorophenyl)-5-(2-methylsulfanylpirimidin-4-yl)-1*H*-pyrazolo[3,4-*b*]pyridine**

Following a similar procedure to that described in reference example 15 section c, but using 2-chloro-6-(4-fluorophenyl)-5-(2-methylsulfanylpirimidin-4-yl)pyridine-3-carbonitrile (obtained in reference example 22) instead of 3-(1-benzylpiperidin-4-yl)-3-oxopropiononitrile, the title compound was obtained.

15

LC-MS (method 1): t_R = 6.70 min; m/z = 353.0 $[\text{M}+\text{H}]^+$.

EXAMPLE 181**6-(4-Fluorophenyl)-5-(2-methylsulfanylpirimidin-4-yl)-1*H*-pyrazolo[3,4-*b*]pyridine**

20

To a solution of 3-amino-6-(4-fluorophenyl)-5-(2-methylsulfanylpirimidin-4-yl)-1*H*-pyrazolo[3,4-*b*]pyridine (10.00 g, 28.4 mmol, obtained in example 180) in AcOH (52 mL), water (22 mL) and HCl conc. (5.7 mL), cooled to 0 °C, a solution of NaNO_2 (2.30 g, 33.4 mmol) in water (7.5 mL) was added dropwise. It was stirred for 30 min at 0 °C, and H_3PO_2 (50% aqueous solution, 56.8 mL) was added slowly. It was stirred at 0 °C for 6 h. It was allowed to cool to room temperature, basified at 0 °C by slow addition of 6 N NaOH hasta pH = 8 and was extracted with EtOAc. The combined organic phases were dried over Na_2SO_4 and concentrated to dryness. The crude product obtained was purified by chromatography on silica gel using hexane-EtOAc mixtures of increasing polarity as eluent, to afford 4.00 g of the title compound (yield: 42%)

25

30

LC-MS (method 1): t_R = 7.80 min; m/z = 338.0 $[\text{M}+\text{H}]^+$.

EXAMPLES 182-193

Following a similar procedure to that described in examples 6 and 7, but starting from the appropriate compounds in each case, the compounds in the following table were obtained:

Example	Compound name	Starting compounds	LC-MS		
			Method	t _R (min)	m/z [M+H] ⁺
182	4,6-Diphenyl-5-(4-pyridyl)-2-[2-(tetrahydropyran-2-yloxy)ethyl]-pyrazolo[3,4-b]pyridine	Example 2 and 2-(2-bromoethoxy)tetrahydropyran	1	6.70	477.1
183	6-(4-Fluorophenyl)-4-(2-furyl)-2-methyl-5-(4-pyridyl)pyrazolo[3,4-b]pyridine	Example 171 and iodomethane	1	5.32	371.1
184	6-(4-Fluorophenyl)-2-methyl-4-(1-methyl-1H-imidazol-4-yl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridine	Example 172 and iodomethane (2 equivalents)	1	3.66	385.2
185	6-(4-Fluorophenyl)-5-(2-methylsulfanylpurimidin-4-yl)-2-[2-(tetrahydropyran-2-yloxy)ethyl]-pyrazolo[3,4-b]pyridine	Example 181 and 2-(2-bromoethoxy)-tetrahydropyran	1	8.79	466.1
186	6-(4-Fluorophenyl)-5-(2-methylsulfanylpurimidin-4-yl)-2-[3-(tetrahydropyran-2-yloxy)propyl]-pyrazolo[3,4-b]pyridine	Example 181 and 2-(3-bromopropoxy)-tetrahydropyran	1	9.11	480.2
187	4,6-Bis(4-fluorophenyl)-5-(2-methylsulfanylpurimidin-4-yl)-2-[3-(tetrahydropyran-2-yloxy)propyl]-pyrazolo[3,4-b]pyridine	Example 174 and 2-(3-bromopropoxy)-tetrahydropyran	1	10.26	574.2
188	4-(5-Bromothien-2-yl)-6-(4-fluorophenyl)-2-methyl-5-(4-pyridyl)-1H-pyrazolo[3,4-b]pyridine	Example 173 and iodomethane	1	6.94	464.9 466.9
189	6-(4-Fluorophenyl)-2-methyl-5-(2-methylsulfanylpurimidin-4-	Example 181 and iodomethane	1	7.29	352.0

	yl)pyrazolo[3,4- <i>b</i>]pyridine				
190	5-(2-Chloropyridin-4-yl)-4,6-bis(4-fluorophenyl)-2-methylpyrazolo[3,4- <i>b</i>]pyridine	Example 175 and iodomethane	1	8.55	433.0 435.0
191	6-(4-Fluorophenyl)-4-(2-phenylethyl)-5-(4-pyridyl)-2-[3-(tetrahydropyran-2-yloxy)propyl]-pyrazolo[3,4- <i>b</i>]pyridine	Example 176 and 2-(3-bromopropoxy)-tetrahydropyran	1	7.74	537.0
192	4-(6-Chloropyridin-3-yl)-6-(4-fluorophenyl)-5-(4-pyridyl)-2-[3-(tetrahydropyran-2-yloxy)propyl]-pyrazolo[3,4- <i>b</i>]pyridine	Example 177 and 2-(3-bromopropoxy)-tetrahydropyran	1	9.05	544.2 546.2
193	4-(6-Chloropyridin-3-yl)-6-(4-fluorophenyl)-2-methyl-5-(4-pyridyl)pyrazolo[3,4- <i>b</i>]pyridine	Example 177 and iodomethane	1	5.34	416.1 418.1

EXAMPLE 194

5 **5-(2-Methylsulfanylpirimidin-4-yl)-2-[3-(tetrahydropyran-2-yloxy)propyl]-6-(3-trifluoromethylphenyl)pyrazolo[3,4-*b*]pyridine**

Following a similar procedure to that described in examples 6 and 7, but starting from 5-[2-(methylsulfanyl)pyrimidin-4-yl]-6-[3-(trifluoromethyl)phenyl]-1*H*-pyrazolo[3,4-*b*]pyridine (obtained in example 126) and 2-(3-bromopropoxy)tetrahydropyran, the title compound was obtained.

10 ¹H NMR (300 MHz, CDCl₃) δ (TMS): 1.50 - 1.90 (m, 4 H), 2.40 (complex signal, 5 H), 3.37 - 3.50 (m, 4 H), 3.85 (m, 2 H), 4.53 (m, 1 H), 4.65 (m, 2 H), 6.68 (d, J = 5.1 Hz, 1 H), 7.40 (t, J = 7.8 Hz, 1 H), 7.52 (d, J = 7.6 Hz, 1 H), 7.61 (d, J = 7.5 Hz, 1 H), 7.93 (s, 1 H), 8.13 (s, 1 H), 8.32 (d, J = 5.1 Hz, 1 H), 8.49 (s, 1 H).

EXAMPLE 195

15 **6-(4-Fluorophenyl)-2-methyl-5-(4-pyridyl)-4-[5-(3-pyridyl)thien-2-yl]pyrazolo[3,4-*b*]pyridine**

A suspension of 4-(5-bromothiophen-2-yl)-6-(4-fluorophenyl)-2-methyl-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine (0.10 g, 0.2 mmol, obtained in example 188), 3-pyridylboronic acid (0.04 g, 0.3 mmol), K₂CO₃ (0.06 g, 0.4 mmol), Pd(PPh₃)₄ (0.017 g, 0.01 mmol), 1,2-dimethoxyethane (1.31 mL) and water (0.04 mL) was

20

heated at 80 °C under argon atmosphere overnight. It was allowed to cool and diluted with CHCl_3 and water. The combined organic phases were dried over Na_2SO_4 and concentrated to dryness. The crude product obtained was purified by chromatography on silica gel using hexane-EtOAc mixtures of increasing polarity as eluent, to afford 35 mg of the desired compound (yield: 50%).

LC-MS (method 1): $t_R = 5.42$ min; $m/z = 464.0$ $[\text{M}+\text{H}]^+$.

EXAMPLES 196-202

Following a similar procedure to that described in example 91 but using the adequate starting compounds in each case, the compounds in the following table were obtained:

Example	Compound name	Starting compound	LC-MS		
			Method	t_R (min)	m/z $[\text{M}+\text{H}]^+$
196	2-[4,6-Diphenyl-5-(4-pyridyl)pyrazolo-[3,4- <i>b</i>]pyridin-2-yl]ethanol	Example 182	1	4.67	393.0
197	3-[5-(2-Methylsulfanylpirimidin-4-yl)-6-(3-trifluoromethylphenyl)pyrazolo-[3,4- <i>b</i>]pyridin-2-yl]propan-1-ol	Example 194	1	7.67	446.0
198	2-[6-(4-Fluorophenyl)-5-(2-methylsulfanylpirimidin-4-yl)pyrazolo[3,4- <i>b</i>]pyridin-2-yl]ethanol	Example 185	1	6.58	382.0
199	3-[6-(4-Fluorophenyl)-5-(2-methylsulfanylpirimidin-4-yl)pyrazolo[3,4- <i>b</i>]pyridin-2-yl]propan-1-ol	Example 186	1	6.79	396.1
200	3-[4,6-Bis(4-fluorophenyl)-5-(2-methylsulfanylpirimidin-4-yl)pyrazolo[3,4- <i>b</i>]pyridin-2-yl]propan-1-ol	Example 187	1	8.10	490.2
201	3-[6-(4-Fluorophenyl)-4-(2-phenylethyl)-5-(4-pyridyl)pyrazolo[3,4- <i>b</i>]pyridin-2-yl]propan-1-ol	Example 191	1	5.80	453.2
202	3-[4-(6-Chloropyridin-3-yl)-6-(4-fluorophenyl)-5-(4-pyridyl)pyrazolo-[3,4- <i>b</i>]pyridin-2-yl]propan-1-ol	Example 192	1	5.88	460.2 462.2

EXAMPLES 203-207

Following a similar procedure to that described in example 56 but using the adequate starting compounds in each case, the compounds in the following table were obtained:

Example	Compound name	Starting compound	LC-MS		
			Method	t _R (min)	m/z [M+H] ⁺
203	6-(4-Fluorophenyl)-2-methyl-5-(2-methylsulfonylpyrimidin-4-yl)pyrazolo[3,4-b]pyridine	Example 189	1	5.78	384.0
204	3-[5-(2-Methylsulfonylpyrimidin-4-yl)-6-(3-trifluoromethylphenyl)pyrazolo-[3,4-b]pyridin-2-yl]propan-1-ol	Example 197	1	6.39	478.0
205	2-[6-(4-Fluorophenyl)-5-(2-methylsulfonylpyrimidin-4-yl)pyrazolo[3,4-b]pyridin-2-yl]ethanol	Example 198	1	5.28	414.0
206	3-[6-(4-Fluorophenyl)-5-(2-methylsulfonylpyrimidin-4-yl)pyrazolo[3,4-b]pyridin-2-yl]propan-1-ol	Example 199	1	5.46	428.0
207	3-[4,6-Bis(4-fluorophenyl)-5-(2-methylsulfonylpyrimidin-4-yl)pyrazolo[3,4-b]pyridin-2-yl]propan-1-ol	Example 200	1	6.76	522.2

5

EXAMPLE 208

N-Cyclopropylmethyl-[4-[6-[3-(trifluoromethyl)phenyl]-1H-pyrazolo[3,4-b]pyridin-5-yl]pyrimidin-2-yl]amine

A solution of 5-[2-(methylsulfonyl)pyrimidin-4-yl]-6-[3-(trifluoromethyl)phenyl]-1H-pyrazolo[3,4-b]pyridine (90 mg, 0.2 mmol, obtained in example 127) and (cyclopropylmethyl)amine (75 mg, 1.0 mmol) in THF (2 mL) was heated in a closed vessel at 60 °C overnight. It was allowed to cool and concentrated. The crude product obtained was purified by chromatography on silica gel using hexane-EtOAc mixtures of increasing polarity as eluent, to afford 57 mg of the desired compound (yield: 67%).

15 LC-MS (method 1): t_R = 8.12 min; m/z = 411.0 [M+H]⁺.

Following a similar procedure to that described in example 208 but using the adequate starting compounds in each case, the compounds in the following table were obtained:

Example	Compound name	Starting compounds	LC-MS		
			Method	t _R (min)	m/z [M+H] ⁺
209	(1S)-3-[5-[2-(1-Phenylethylamino)pyrimidin-4-yl]-6-(3-trifluoromethylphenyl)pyrazolo[3,4-b]pyridin-2-yl]propan-1-ol	Example 204 and (1S)-1-phenylethylamine	1	9.11	519.1
210	N-Cyclopropylmethyl-[4-[6-(4-fluorophenyl)-2-methylpyrazolo[3,4-b]pyridin-5-yl]pyrimidin-2-yl]amine	Example 203 and (cyclopropylmethyl)amine	1	6.43	375.1
211	2-[5-[2-[(Cyclopropylmethyl)amino]pyrimidin-4-yl]-6-(4-fluorophenyl)pyrazolo[3,4-b]pyridin-2-yl]ethanol	Example 205 and (cyclopropylmethyl)amine	1	5.85	405.1
212	3-[5-[2-[(Cyclopropylmethyl)amino]pyrimidin-4-yl]-6-(4-fluorophenyl)pyrazolo[3,4-b]pyridin-2-yl]propan-1-ol	Example 206 and (cyclopropylmethyl)amine	1	6.06	419.1
213	3-[5-[2-[(Cyclopropylmethyl)amino]pyrimidin-4-yl]-4,6-bis(4-fluorophenyl)pyrazolo[3,4-b]pyridin-2-yl]propan-1-ol	Example 207 and (cyclopropylmethyl)amine	1	7.63	513.3

5

EXAMPLE 214

4-[4-[4,6-Bis(4-fluorophenyl)-2-methylpyrazolo[3,4-b]pyridin-5-yl]pyridin-2-ylamino]benzenesulfonamide

A mixture of 5-(2-chloropyridin-4-yl)-4,6-bis(4-fluorophenyl)-2-methylpyrazolo[3,4-b]pyridine (100 mg, 0.23 mmol, obtained in example 190) and 4-aminobenzenesulfonamide (46 mg, 0.27 mmol) was heated at 190 °C overnight. It was allowed to cool and the crude product obtained was purified by

10

chromatography on silica gel using EtOAc as eluent, to afford 23 mg of the desired compound (yield: 17%).

LC-MS (method 1): t_R = 6.91 min; m/z = 569.0 $[M+H]^+$.

EXAMPLES 215-230

- 5 Following a similar procedure to that described in example 72 but using the adequate starting compounds in each case, the compounds in the following tables were obtained:

Example	Compound name	Starting compounds	LC-MS		
			Method	t_R (min)	m/z $[M+H]^+$
215	4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo-[3,4-b]pyridin-3-ol	Reference example 1, 5-amino-1 <i>H</i> -pyrazol-3-ol and 4-fluorobenzaldehyde	1	5.02	401.1
216	6-(4-Fluorophenyl)-4-(3 <i>H</i> -imidazol-4-yl)-5-(4-pyridyl)-1 <i>H</i> -pyrazolo[3,4-b]pyridine	Reference example 1, 3-amino-2 <i>H</i> -pyrazole and 3 <i>H</i> -imidazol-4-carbaldehyde	1	3.23	357.1
217	6-(4-Fluorophenyl)-4-(1 <i>H</i> -pyrazol-3-yl)-5-(4-pyridyl)-1 <i>H</i> -pyrazolo[3,4-b]pyridine	Reference example 1, 3-amino-2 <i>H</i> -pyrazole and 2 <i>H</i> -pyrazole-3-carbaldehyde	1	4.24	357.1
218	3-[6-(4-Fluorophenyl)-5-(4-pyridyl)-1 <i>H</i> -pyrazolo-[3,4-b]pyridin-4-yl]phenol	Reference example 1, 3-amino-2 <i>H</i> -pyrazole and 3-hydroxybenzaldehyde	1	4.99	383.1
219	4-Cyclopropyl-6-(4-fluorophenyl)-5-(4-pyridyl)-1 <i>H</i> -pyrazolo[3,4-b]pyridine	Reference example 1, 3-amino-2 <i>H</i> -pyrazole and cyclopropanecarbaldehyde	1	4.99	331.1
220	6-(4-Fluorophenyl)-4-(5-methylfuran-2-yl)-5-(4-pyridyl)-1 <i>H</i> -pyrazolo[3,4-b]pyridine	Reference example 1, 3-amino-2 <i>H</i> -pyrazole and 5-methylfuran-2-carbaldehyde	1	8.45	445
221	4-(5-Bromofuran-2-yl)-6-(4-fluorophenyl)-5-(4-pyridyl)-1 <i>H</i> -pyrazolo[3,4-	Reference example 1, 3-amino-2 <i>H</i> -pyrazole and 5-methylfuran-2-carbaldehyde	1	7.02	435.0 437.0

	b]pyridine				
222	4-(4-Benzoyloxyphenyl)-6-(4-fluorophenyl)-2-methyl-5-pyrimidin-4-yl-1H-pyrazolo[3,4-b]pyridine	Reference example 30, 3-amino-2H-pyrazole and 4-benzoyloxybenzaldehyde	1	9.05	474.1
223	4-(4-Benzoyloxyphenyl)-6-(4-fluorophenyl)-5-(2-methylsulfanylpymidin-4-yl)-1H-pyrazolo[3,4-b]pyridine	Reference example 19, 3-amino-2H-pyrazole and 4-benzoyloxybenzaldehyde	1	10.52	520.1
224	6-(4-Fluorophenyl)-4-propyl-5-(4-pyridyl)-1H-pyrazolo[3,4-b]pyridine	Reference example 1, 3-amino-2H-pyrazole and butyraldehyde	1	5.42	333.0
225	4-(3-Benzoyloxyphenyl)-6-(4-fluorophenyl)-5-(4-pyridyl)-1H-pyrazolo[3,4-b]pyridine	Reference example 1, 3-amino-2H-pyrazole and 3-benzoyloxybenzaldehyde	1	8.00	473.2
226	5-(2-Chloropyridin-4-yl)-6-(4-fluorophenyl)-1H-pyrazolo[3,4-b]pyridine	Reference example 23, 3-amino-2H-pyrazole and paraformaldehyde	1	7.73	325.2 327.2
227	4-[6-(4-Fluorophenyl)-5-(4-pyridyl)-1H-pyrazolo[3,4-b]pyridin-4-yl]butan-1-ol	Reference example 1, 3-amino-2H-pyrazole and 5-hydroxypentanal	1	4.23	363.2
228	4-Benzyl-6-(4-fluorophenyl)-5-(4-pyridyl)-1H-pyrazolo[3,4-b]pyridine	Reference example 1, 3-amino-2H-pyrazole and phenylacetaldehyde	1	6.37	381.2

Example	Compound name	Starting compounds	¹ H NMR (300 MHz, CDCl ₃) δ (TMS)
229	4-(4-Benzoyloxyphenyl)-6-(4-fluorophenyl)-2-methyl-5-(4-pyridyl)-1H-pyrazolo[3,4-b]pyridine	Reference example 1, 3-amino-2H-pyrazole and 4-benzyloxy-benzaldehyde	5.06 (s, 2 H), 6.85 (dd, J _o = 1.5 Hz, J _m = 4.5 Hz, 2 H), 6.91 (m, 4 H), 6.95 (m, 2 H), 7.23 - 7.40 (m, 7 H), 8.01 (s, 1 H), 8.35 (dd, J _o = 1.5 Hz, J _m = 4.5 Hz, 2 H), 10.62 (broad s, 1 H, NH).
230	4,6-Bis(4-fluorophenyl)-5-pyrimidin-4-yl-1H-pyrazolo[3,4-b]pyridine	Reference example 30, 3-amino-2H-pyrazole and 4-fluorobenzaldehyde	6.92 - 7.08 (complex signal, 5 H), 7.23 (m, 2 H), 7.32 (m, 2 H), 7.99 (s, 1 H), 8.45 (d, J = 5.1 Hz, 1 H), 9.03 (s, 1 H), 11.40 (broad s, 1 H, NH).

EXAMPLE 231

[(2S)-2-[4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridin-2-yl]ethyl]pyrrolidine-2-carboxamide

- 5 a) [(2S)-2-[4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridin-2-yl]ethyl]-1-(benzyloxycarbonyl)pyrrolidine-2-carboxamide

Following a similar procedure to that described in example 48, but starting from 2-(2-aminoethyl)-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridine (obtained in example 90) and (2S)-1-(benzyloxycarbonyl)pyrrolidine-2-carboxylic acid, the desired compound was obtained.

LC-MS (method 1): t_R = 7.13 min; m/z = 659.3 [M+H]⁺.

b) **Title compound**

Following a similar procedure to that described in example 269, but starting from [2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridin-2-yl]ethyl]-1-(benzyloxycarbonyl)pyrrolidine-2-carboxamide (obtained in section a), the title compound was obtained.

LC-MS (method 1): t_R = 4.35 min; m/z = 525.2 [M+H]⁺.

EXAMPLE 232

2-[2-(4,6-Diphenyl-5-(4-pyridyl)pyrazolo[3,4-b]pyridin-2-yl)ethylamino]ethanol

- 20 a) **2-[4,6-Diphenyl-5-(4-pyridyl)pyrazolo[3,4-b]pyridin-2-yl]ethyl methanesulfonate**

Following a similar procedure to that described in example 12 section b, but starting from 2-[4,6-diphenyl-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethanol (obtained in example 196), the desired compound was obtained.

¹H NMR (300 MHz, CDCl₃) δ (TMS): 2.94 (s, 3 H), 4.76 (m, 2 H), 4.85 (m, 2 H),
 5 6.88 (d, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H), 7.15 - 7.90 (complex signal, 10 H), 8.27 (s, 1H), 8.28 (dd, J_o = 1.5 Hz, J_m = 4.5 Hz, 2 H).

b) Title compound

Following a similar procedure to that described in example 111, but starting from 2-[4,6-diphenyl-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl methanesulfonate
 10 (obtained in section a) and 2-amino-1-ethanol, the desired compound was obtained.

LC-MS (method 1): t_R = 3.89 min; m/z = 436.1 [M+H]⁺.

EXAMPLE 233

6-(4-Fluorophenyl)-2-methyl-4-(3-pyridyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine
 15 To a solution of 4-(6-chloropyridin-3-yl)-6-(4-fluorophenyl)-2-methyl-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine (74 mg, 0.2 mmol, obtained in example 193) in AcOH (1 mL), Zn (72 mg, 1.1 mmol) was added under argon atmosphere and the mixture was heated to reflux overnight. It was allowed to cool and concentrated. The residue was treated with saturated NaHCO₃ and it was extracted with CHCl₃.
 20 The organic phase was dried over Na₂SO₄ and concentrated to dryness. The crude product obtained was purified by chromatography on silica gel using hexane-EtOAc-MeOH mixtures of increasing polarity, to afford 2.4 mg of the title compound (yield: 4%)

LC-MS (method 1): t_R = 4.02 min; m/z = 382.2 [M+H]⁺.

25

EXAMPLES 234-235

Following a similar procedure to that described in reference example 15 section c, but starting from the appropriate compounds in each case, the compounds in the following table were obtained:

Example	Compound name	Starting compounds	LC-MS		
			Method	t _R (min)	m/z [M+H] ⁺
234	6-(4-Fluorophenyl)-3-methyl-5-(2-methylsulfanylpirimidin-	Reference example 29 and hydrazine	1	8.57	352.0

	4-yl)-1 <i>H</i> -pyrazolo[3,4- <i>b</i>]pyridine	monohydrate			
235	3-Amino-5-(2-methylsulfanyl- pyrimidin-4-yl)-6-phenyl-1 <i>H</i> - pyrazolo[3,4- <i>b</i>]pyridine	Reference example 28 and hydrazine monohydrate	1	6.55	335.0

EXAMPLES 236-237

Following a similar procedure to that described in example 77, but starting from the appropriate compounds in each case, the compounds in the following table were obtained:

Example	Compound name	Starting compound	LC-MS		
			Method	t _R (min)	m/z [M+H] ⁺
236	5-(2-Methylsulfanylpymidin-4-yl)- 6-phenyl-1 <i>H</i> -pyrazolo[3,4- <i>b</i>]pyridine	Example 235	1	7.54	320.0
237	6-(4-Fluorophenyl)-4-methyl-5-(2-methylsulfanylpymidin-4-yl)-1 <i>H</i> -pyrazolo[3,4- <i>b</i>]pyridine	Example 147	1	4.42	305.0

EXAMPLE 238

5-(2-Methoxypyrimidin-4-yl)-6-(3-trifluoromethylphenyl)-1*H*-pyrazolo[3,4-*b*]pyridine

- 10 To a solution of 5-[2-(methylsulfonyl)pyrimidin-4-yl]-6-[3-(trifluoromethyl)phenyl]-1*H*-pyrazolo[3,4-*b*]pyridine (90 mg, 0.2 mmol, obtained in example 127) in MeOH (5 mL) in a closed vessel, sodium methoxide (11 mg, 0.2 mmol) was added and heated to 60 °C for 24 h. Then, sodium methoxide (11 mg, 0.2 mmol) was added and it was stirred at 60 °C for 2 h more. It was allowed to cool and was
- 15 concentrated. EtOAc and buffer (pH = 5.3) were added. The phases were separated and the organic phase was washed with water, dried over Na₂SO₄ and concentrated to dryness. The crude product obtained was purified by chromatography on silica gel using hexane-EtOAc mixtures at 50%, to afford 38 mg of the title compound (yield: 47%)
- 20 LC-MS (method 1): t_R = 7.71 min; m/z = 372.0 [M+H]⁺.

EXAMPLES 239-242

Following a similar procedure to that described in reference example 1 section a, but starting from the appropriate compounds in each case, the compounds in the following table were obtained:

Example	Compound name	Starting compound	LC-MS		
			Method	t _R (min)	m/z [M+H] ⁺
239	N-[2-[4,6-(Diphenyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridin-2-yl]ethyl]acetamide	Example 142 and acetyl chloride	1	5.34	434.1
240	N-[3-[4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridin-2-yl]propyl]acetamide	Example 95 and acetyl chloride	1	5.31	484.1
241	N-[2-(4,6-Diphenyl-5-(4-pyridyl)pyrazolo[3,4-b]pyridin-2-yl)ethyl]-N-(2-hydroxyethyl)-acetamide	Example 232 and acetyl chloride	1	5.27	478.1
242	N-[2-[4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridin-2-yl]ethyl]propionamide	Example 90 and propionyl chloride	1	5.39	484.1

5

EXAMPLE 243

N-[3-[4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridin-2-yl]propyl]methanesulfonamide

Following a similar procedure to that described in example 169, but starting from 2-(3-aminopropyl)-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridine (obtained in example 95), the desired compound was obtained.

10

LC-MS (method 1): t_R = 5.79 min; m/z = 520.1 [M+H]⁺.

EXAMPLE 244

5-(2-Aminopyrimidin-4-yl)-6-(3-trifluoromethylphenyl)-1H-pyrazolo[3,4-b]pyridine

15 A solution of THF (20 mL) saturated with NH₃ (g) at -20 °C was added over 5-[2-(methylsulfonyl)pyrimidin-4-yl]-6-[3-(trifluoromethyl)phenyl]-1H-pyrazolo[3,4-b]pyridine (90 mg, 0.2 mmol, obtained in example 127) in a closed vessel. It was stirred at room temperature for 2 days and concentrated. The crude product obtained was purified by chromatography on silica gel using hexane-EtOAc

mixtures of increasing polarity as eluent, to afford 7 mg of the title compound (yield: 9%)

LC-MS (method 1): t_R = 6.01 min; m/z = 357.0 $[M+H]^+$.

EXAMPLE 245

5 ***N*-[5-(2-Methylsulfanylpirimidin-4-yl)-6-(3-trifluoromethylphenyl)-1*H*-pyrazolo[3,4-*b*]pyridin-3-yl]acetamide**

A solution of 3-amino-5-[2-(methylsulfanyl)pyrimidin-4-yl]-6-[3-(trifluoromethyl)phenyl]-1*H*-pyrazolo[3,4-*b*]pyridine (200 mg, 0.5 mmol, obtained in example 123) and acetyl chloride (39 mg, 0.5 mmol) in pyridine (10 mL) was
10 stirred at room temperature for 3 h. It was concentrated and the residue was taken up in a mixture of EtOAc and 1 N NaOH. The organic phase was dried over Na₂SO₄ and concentrated to dryness. The crude product obtained was purified by chromatography on silica gel using hexane-EtOAc mixtures of increasing polarity as eluent, to afford 78 mg of the title compound (yield: 37%)

15 LC-MS (method 1): t_R = 8.55 min; m/z = 445.0 $[M+H]^+$.

EXAMPLE 246

***N*-Cyclopropylmethyl-[4-[3-benzyloxycarbonylamino-6-(3-trifluoromethylphenyl)-1*H*-pyrazolo[3,4-*b*]pyridin-5-yl]pyrimidin-2-yl]amine**
a) ***N*-Benzyloxycarbonyl-[5-(2-methylsulfanylpirimidin-4-yl)-6-(3-trifluoromethylphenyl)-1*H*-pyrazolo[3,4-*b*]pyridin-3-yl]amine**
20

Following a similar procedure to that described in example 245, but starting from 3-amino-5-[2-(methylsulfanyl)pyrimidin-4-yl]-6-[3-(trifluoromethyl)phenyl]-1*H*-pyrazolo[3,4-*b*]pyridine (obtained in example 123) and benzyl chloroformate, the desired compound was obtained.

25 LC-MS (method 1): t_R = 10.00 min; m/z = 537.1 $[M+H]^+$.

b) ***N*-Benzyloxycarbonyl-[5-(2-methylsulfonylpirimidin-4-yl)-6-(3-trifluoromethylphenyl)-1*H*-pyrazolo[3,4-*b*]pyridin-3-yl]amine**

Following a similar procedure to that described in example 56, but starting from *N*-benzyloxycarbonyl-[5-(2-methylsulfanylpirimidin-4-yl)-6-(3-trifluoromethylphenyl)-1*H*-pyrazolo[3,4-*b*]pyridin-3-yl]amine (obtained in section a), the desired
30 compound was obtained.

LC-MS (method 1): t_R = 8.47 min; m/z = 567.1 $[M+H]^+$.

c) Title compound

Following a similar procedure to that described in example 208, but starting from *N*-benzyloxycarbonyl-[5-(2-methylsulfonylpyrimidin-4-yl)-6-(3-trifluoromethylphenyl)-1*H*-pyrazolo[3,4-*b*]pyridin-3-yl]amine (obtained in section b) and (cyclopropylmethyl)amine, the desired compound was obtained.

5 LC-MS (method 1): $t_R = 9.44$ min; $m/z = 560.3$ $[M+H]^+$.

EXAMPLE 247

***N*-[2-[4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl]-2-hydroxyacetamide**

Following a similar procedure to that described in example 48, but starting from 2-
10 (2-aminoethyl)-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine (obtained in example 90) and hydroxyacetic acid, the title compound was obtained.

LC-MS (method 1): $t_R = 4.84$ min; $m/z = 486.1$ $[M+H]^+$.

EXAMPLE 248

15 ***N*-[2-[4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl]piperidine-4-carboxamide**

a) ***N*-[2-[4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl]-1-(*tert*-butoxycarbonyl)piperidine-4-carboxamide**

Following a similar procedure to that described in example 48, but starting from 2-
20 (2-aminoethyl)-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine (obtained in example 90) and 1-(*tert*-butoxycarbonyl)piperidine-4-carboxylic acid, the desired compound was obtained.

b) **Title compound**

Following a similar procedure to that described in example 36, but starting from *N*-
25 [2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl]-1-(*tert*-butoxycarbonyl)piperidine-4-carboxamide (obtained in section a), the title compound was obtained.

LC-MS (method 1): $t_R = 4.29$ min; $m/z = 539.2$ $[M+H]^+$.

EXAMPLE 249

30 ***N*-[2-[4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl]-2-(methylamino)acetamide**

a) ***N*-[2-[4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl]-2-chloroacetamide**

Following a similar procedure to that described in reference example 1 section a, but starting from 2-(2-aminoethyl)-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine (obtained in example 90) and chloroacetyl chloride, the title compound was obtained.

5 LC-MS (method 1): t_R = 5.72 min; m/z = 504.1, 506.1 $[M+H]^+$.

b) Title compound

Following a similar procedure to that described in example 78, but starting from *N*-[2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl]-2-chloroacetamide (obtained in section a) and methylamine, the title compound was

10 obtained.

LC-MS (method 1): t_R = 4.28 min; m/z = 499.2 $[M+H]^+$.

EXAMPLE 250

***N*-[2-[4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl]-2-(2-hydroxyethylamino)acetamide**

15 Following a similar procedure to that described in example 78, but starting from *N*-[2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl]-2-chloroacetamide (obtained in example 249 section a) and 2-aminoethanol, the title compound was obtained.

LC-MS (method 1): t_R = 4.29 min; m/z = 529.2 $[M+H]^+$.

20

EXAMPLE 251

***N*-[2-[4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl]nicotinamide**

a) Nicotinoyl chloride hydrochloride

25 A solution of nicotinic acid (0.50 g, 4.0 mmol) and POCl₃ (5 mL) was heated at reflux for 1.5 h and concentrated. The product obtained was directly used in the next reaction.

b) Title compound

Following a similar procedure to that described in example 245, but starting from 2-(2-aminoethyl)-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine

30 (obtained in example 90) and nicotinoyl chloride hydrochloride (obtained in section a), the title compound was obtained.

LC-MS (method 1): t_R = 5.25 min; m/z = 533.1 $[M+H]^+$.

EXAMPLES 252-265

Following a similar procedure to that described in examples 6 and 7, but starting from the appropriate compounds in each case, the compounds in the following table were obtained:

Example	Compound name	Starting compound	LC-MS		
			Method	t _R (min)	m/z [M+H] ⁺
252	4-(4-Benzoyloxyphenyl)-6-(4-fluorophenyl)-2-methyl-5-(4-pyridyl)-pyrazolo[3,4-b]pyridine	Example 229 and iodomethane	1	7.53	487.1
253	6-(4-Fluorophenyl)-5-(2-methylsulfanylpirimidin-4-yl)-2-methyl-pyrazolo[3,4-b]pyridine	Example 181 and iodomethane	1	7.13	352.1
254	6-(4-Fluorophenyl)-2,4-dimethyl-5-(4-pyridyl)pyrazolo[3,4-b]pyridine	Example 237 and iodomethane	1	4.11	319.00
255	4-(4-Benzoyloxyphenyl)-6-(4-fluorophenyl)-2-methyl-5-(2-methylsulfanylpirimidin-4-yl)pyrazolo[3,4-b]pyridine	Example 223 and iodomethane	1	10.16	534.2
256	2-(1-Benzylpyrrolidin-2-ylmethyl)-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)-pyrazolo[3,4-b]pyridine	Example 1 and 1-benzyl-2-chloromethylpyrrolidine	1	6.10	558.2
257	4-(4-Benzoyloxyphenyl)-6-(4-fluorophenyl)-2-methyl-5-pyrimidin-4-ylpyrazolo[3,4-b]pyridine	Example 222 and iodomethane	1	8.72	488.2
258	6-(4-Fluorophenyl)-2-methyl-4-(5-methylfuran-2-yl)-5-(4-pyridyl)-pyrazolo[3,4-b]pyridine	Example 220 and iodomethane	1	5.85	385.0
259	4-(5-Bromofuran-2-yl)-6-(4-fluorophenyl)-2-methyl-5-(4-pyridyl)-pyrazolo[3,4-b]pyridine	Example 221 and iodomethane	1	6.48	449.1 451.1
260	6-(4-Fluorophenyl)-2-methyl-4-propyl-5-(4-pyridyl)pyrazolo[3,4-b]pyridine	Example 224 and iodomethane	1	4.88	347.0

261	4-(3-Benzoyloxyphenyl)-6-(4-fluorophenyl)-2-methyl-5-(4-pyridyl)-pyrazolo[3,4- <i>b</i>]pyridine	Example 225 and iodomethane	1	7.79	487.0
262	6-(4-Fluorophenyl)-4-(2-phenyl)ethyl-2-methyl-5-(4-pyridyl)pyrazolo[3,4- <i>b</i>]pyridine	Example 176 and iodomethane	1	5.84	409.0
263	5-(2-Chloropyridin-4-yl)-6-(4-fluorophenyl)-2-methylpyrazolo[3,4- <i>b</i>]pyridine	Example 228 and iodomethane	1	7.30	339.2 341.2
264	4-Benzyl-6-(4-fluorophenyl)-2-methyl-5-(4-pyridyl)pyrazolo[3,4- <i>b</i>]pyridine	Example 228 and iodomethane	1	5.95	395.2
265	4-[6-(4-Fluorophenyl)-2-methyl-5-(4-pyridyl)pyrazolo[3,4- <i>b</i>]pyridin-4-yl]butan-1-ol	Example 227 and iodomethane	1	3.69	377.2

EXAMPLE 266**4-[6-(4-Fluorophenyl)-2-methyl-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-4-yl]phenol**

5

To a solution of 4-(4-benzoyloxyphenyl)-6-(4-fluorophenyl)-2-methyl-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine (112 mg, 0.2 mmol, obtained in example 252) in EtOH (13 mL), 10% Pd/C (20 mg) was added and it was hydrogenated under atmospheric pressure at room temperature for 2 days. It was filtered through celite, washed with EtOH and concentrated. The crude product obtained was purified by chromatography on silica gel using hexane-EtOAc-MeOH mixtures of increasing polarity as eluent, to afford 61 mg of the title compound (yield: 67%)

LC-MS (method 1): t_R = 4.65 min; m/z = 397.1 $[M+H]^+$.

10

EXAMPLE 267***N*-[6-(4-Fluorophenyl)-5-(2-methylsulfanylpirimidin-4-yl)-1*H*-pyrazolo[3,4-*b*]pyridin-3-yl]acetamide**

15

Following a similar procedure to that described in example 245, but starting from 3-amino-6-(4-fluorophenyl)-5-(2-methylsulfanylpirimidin-4-yl)pyrazolo[3,4-*b*]pyridine (obtained in example 180) and acetyl chloride, the title compound was obtained.

20

LC-MS (method 1): $t_R = 7.01$ min; $m/z = 395.1$ $[M+H]^+$.

EXAMPLE 268

***N*-[5-[2-[(Cyclopropylmethyl)amino]pyrimidin-4-yl]-6-(4-fluorophenyl)-1*H*-pyrazolo[3,4-*b*]pyridin-3-yl]acetamide**

- 5 a) ***N*-[6-(4-Fluorophenyl)-5-(2-methylsulfonylpyrimidin-4-yl)-1*H*-pyrazolo[3,4-*b*]pyridin-3-yl]acetamide** and ***N*-[6-(4-fluorophenyl)-5-(2-methylsulfonylpyrimidin-4-yl)-1*H*-pyrazolo[3,4-*b*]pyridin-3-yl]acetamide 7-oxide**

10 Following a similar procedure to that described in example 56, but starting from *N*-[6-(4-fluorophenyl)-5-(2-methylsulfonylpyrimidin-4-yl)-1*H*-pyrazolo[3,4-*b*]pyridin-3-yl]acetamide (obtained in example 267) and using 2 equivalents of *m*-chloroperbenzoic acid, the title compounds were obtained.

N-[6-(4-Fluorophenyl)-5-(2-methylsulfonylpyrimidin-4-yl)-1*H*-pyrazolo[3,4-*b*]pyridin-3-yl]acetamide: LC-MS (method 1): $t_R = 5.64$ min; $m/z = 427.1$ $[M+H]^+$.

- 15 *N*-[6-(4-Fluorophenyl)-5-(2-methylsulfonylpyrimidin-4-yl)-1*H*-pyrazolo[3,4-*b*]pyridin-3-yl]acetamide 7-oxide: LC-MS (method 1): $t_R = 4.44$ min; $m/z = 443.0$ $[M+H]^+$.

b) Title compound

20 Following a similar procedure to that described in example 208, but starting from *N*-[6-(4-fluorophenyl)-5-(2-methylsulfonylpyrimidin-4-yl)-1*H*-pyrazolo[3,4-*b*]pyridin-3-yl]acetamide (obtained in section a) and (cyclopropylmethyl)amine, the title compound was obtained.

LC-MS (method 1): $t_R = 6.19$ min; $m/z = 418.1$ $[M+H]^+$.

EXAMPLE 269

25 **3-[6-(4-Fluorophenyl)-4-(2-furyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]propan-1-ol**

- a) **6-(4-Fluorophenyl)-4-(2-furyl)-5-(4-pyridyl)-2-[3-(tetrahydropyran-2-yloxy)propyl]pyrazolo[3,4-*b*]pyridine**

30 Following a similar procedure to that described in examples 6 and 7, but starting from 6-(4-fluorophenyl)-4-(2-furyl)-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine (obtained in example 171) and 2-(3-bromopropoxy)tetrahydropyran, the desired compound was obtained.

LC-MS (method 1): $t_R = 7.19$ min; $m/z = 499.2$ $[M+H]^+$.

b) Title compound

Following a similar procedure to that described in example 91, but starting from 6-(4-fluorophenyl)-4-(2-furyl)-5-(4-pyridyl)-2-[3-(tetrahydropyran-2-yloxy)propyl]pyrazolo[3,4-*b*]pyridine (obtained in section a), the title compound was obtained.

- 5 LC-MS (method 1): $t_R = 5.05$ min; $m/z = 415.1$ $[M+H]^+$.

Following a similar procedure to that described in example 269, but starting from the appropriate compounds in each case, the compounds in the following table were obtained:

Example	Compound name	Starting compound	LC-MS		
			Method	t_R (min)	m/z $[M+H]^+$
270	2-[4-(4-Benzoyloxyphenyl)-6-(4-fluorophenyl)-5-(4-pyridyl)-pyrazolo[3,4- <i>b</i>]pyridin-2-yl]ethanol	Example 229 and 2-(2-bromoethoxy)-tetrahydropyran	1	6.76	517.2
271	3-[4-(4-Benzoyloxyphenyl)-6-(4-fluorophenyl)-5-(4-pyridyl)-pyrazolo[3,4- <i>b</i>]pyridin-2-yl]propan-1-ol	Example 229 and 2-(3-bromopropoxy)-tetrahydropyran	1	6.91	531.2
272	3-[6-(4-Fluorophenyl)-4-(5-methylfuran-2-yl)-5-(4-pyridyl)-pyrazolo[3,4- <i>b</i>]pyridin-2-yl]propan-1-ol	Example 220 and 2-(3-bromopropoxy)-tetrahydropyran	1	5.42	429.1
273	3-[4-Cyclopropyl-6-(4-fluorophenyl)-5-(4-pyridyl)-pyrazolo[3,4- <i>b</i>]pyridin-2-yl]propan-1-ol	Example 219 and 2-(3-bromopropoxy)-tetrahydropyran	1	4.40	389.1
274	3-[4-(5-Bromothien-2-yl)-6-(4-fluorophenyl)-5-(4-pyridyl)-pyrazolo[3,4- <i>b</i>]pyridin-2-yl]propan-1-ol	Example 173 and 2-(3-bromopropoxy)-tetrahydropyran	1	6.25	508.9 510.9
275	3-[6-(4-Fluorophenyl)-4-propyl-5-(4-pyridyl)pyrazolo[3,4- <i>b</i>]pyridin-2-yl]propan-1-ol	Example 224 and 2-(3-bromopropoxy)-tetrahydropyran	1	4.90	391.2

276	3-[4,6-Bis(4-fluorophenyl)-5-pyrimidin-4-ylpyrazolo[3,4-b]pyridin-2-yl]propan-1-ol	Example 230 and 2-(3-bromopropoxy)-tetrahydropyran	1	6.56	444.2
277	3-[4-(3-Benzyloxyphenyl)-6-(4-fluorophenyl)-5-(4-pyridyl)-pyrazolo[3,4-b]pyridin-2-yl]propan-1-ol	Example 225 and 2-(3-bromopropoxy)-tetrahydropyran	1	7.03	531.3
278	3-[4-Benzyl-6-(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridin-2-yl]propan-1-ol	Example 228 and 2-(3-bromopropoxy)-tetrahydropyran	1	5.27	439.2

EXAMPLES 279-285

Following a similar procedure to that described in example 268, but starting from the appropriate compound and a suitable amine in each case, the compounds in the following table were obtained:

Example	Compound name	Starting compound	Amine	LC-MS		
				Method	t _R (min)	m/z [M+H] ⁺
279	(1S)-N-(1-Phenylethyl)-4-[6-(4-fluorophenyl)-3-methyl-1H-pyrazolo[3,4-b]pyridin-5-yl]pyrimidin-2-yl]amine	Example 234	(1S)-1-Phenylethylamine	1	9.32	425.1
280	N-Cyclopropylmethyl-4-[6-(4-fluorophenyl)-3-methyl-1H-pyrazolo[3,4-b]pyridin-5-yl]pyrimidin-2-yl]amine	Example 234	(Cyclopropylmethyl)amine	2	5.54	373.2
281	1-[4-[6-(4-Fluorophenyl)-2-methylpyrazolo[3,4-b]pyridin-5-yl]pyrimidin-2-ylamino]propan-2-ol	Example 253	1-Aminopropan-2-ol	1	4.73	379.2
282	N-Cyclopropylmethyl-4-[6-[phenyl-1H-pyrazolo[3,4-b]pyridin-5-yl]pyrimidin-2-yl]amine	Example 236	(Cyclopropylmethyl)amine	1	7.53	343.1
283	2-[4-[6-(4-Fluorophenyl)-2-methylpyrazolo[3,4-	Example 253	2-Aminopropan-	1	4.74	379.1

	b]pyridin-5-yl]pyrimidin-2-ylamino]propan-1-ol		1-ol			
284	4-[4-[6-(4-Fluorophenyl)-2-methylpyrazolo[3,4-b]pyridin-5-yl]pyrimidin-2-ylamino]butan-1-ol	Example 253	4-Aminobutan-1-ol	1	4.61	393.2
285	(1S)-N-(1-Phenylethyl)-4-[6-phenyl-1H-pyrazolo[3,4-b]pyridin-5-yl]pyrimidin-2-ylamine	Example 236	(1S)-1-Phenylethylamine	1	8.10	393.1

EXAMPLES 286-290

Following a similar procedure to that described in example 208, but starting from the appropriate compounds in each case, the compounds in the following table

5 were obtained:

Example	Compound name	Starting compounds	LC-MS		
			Method	t _R (min)	m/z [M+H] ⁺
286	N-(3-Methoxypropyl)-4-[6-(3-trifluoromethylphenyl)-1H-pyrazolo[3,4-b]pyridin-5-yl]pyrimidin-2-ylamine	Example 127 and 3-methoxypropylamine	1	7.14	429.1
287	3-[4-[6-(3-Trifluoromethylphenyl)-1H-pyrazolo[3,4-b]pyridin-5-yl]pyrimidin-2-ylamino]propan-1-ol	Example 127 and 3-amino-propan-1-ol	1	5.91	415.1
288	3-[4-[6-(4-Fluorophenyl)-2-(3-hydroxypropyl)pyrazolo[3,4-b]pyridin-5-yl]pyrimidin-2-ylamino]propan-1-ol	Example 206 and 3-amino-propan-1-ol	1	4.55	423.1
289	N-Ethyl-4-[6-(3-trifluoromethylphenyl)-1H-pyrazolo[3,4-b]pyridin-5-yl]pyrimidin-2-ylamine	Example 127 and ethylamine	1	7.21	385.0
290	N-Benzyl-4-[6-(3-trifluoromethylphenyl)-1H-pyrazolo[3,4-b]pyridin-5-yl]pyrimidin-2-ylamine	Example 127 and benzylamine	1	8.86	447.1

EXAMPLE 291

4-[5-[2-[(Cyclopropylmethyl)amino]pyrimidin-4-yl]-6-(4-fluorophenyl)-2-methylpyrazolo[3,4-b]pyridin-4-yl]phenol

a) 4-(4-Benzoyloxyphenyl)-6-(4-fluorophenyl)-5-(2-methylsulfonylpyrimidin-4-yl)-2-methylpyrazolo[3,4-b]pyridine

- 5 Following a similar procedure to that described in example 56, but starting from 4-(4-benzoyloxyphenyl)-6-(4-fluorophenyl)-2-methyl-5-(2-methylsulfonylpyrimidin-4-yl)pyrazolo[3,4-b]pyridine (obtained in example 255), the desired compound was obtained.

LC-MS (method 1): $t_R = 8.77$ min; $m/z = 566.2$ $[M+H]^+$.

10 **b) 4-[6-(4-Fluorophenyl)-5-(2-methylsulfonylpyrimidin-4-yl)-2-methylpyrazolo[3,4-b]pyridin-4-yl]phenol**

Following a similar procedure to that described in example 266, but starting from 4-(4-benzoyloxyphenyl)-6-(4-fluorophenyl)-2-methyl-5-(2-methylsulfonylpyrimidin-4-yl)pyrazolo[3,4-b]pyridine (obtained in section a), the desired compound was

- 15 obtained.

LC-MS (method 1): $t_R = 6.17$ min; $m/z = 476.1$ $[M+H]^+$.

c) Title compound

Following a similar procedure to that described in example 128, but starting from 4-[6-(4-fluorophenyl)-5-(2-methylsulfonylpyrimidin-4-yl)-2-methylpyrazolo[3,4-

- 20 b]pyridin-4-yl]phenol (obtained in section b) and (cyclopropylmethyl)amine, the title compound was obtained.

LC-MS (method 1): $t_R = 6.62$ min; $m/z = 467.2$ $[M+H]^+$.

EXAMPLES 292-295

- 25 Following a similar procedure to that described in example 266, but starting from the appropriate compounds in each case, the compounds in the following table were obtained:

Example	Compound name	Starting compound	LC-MS		
			Method	t_R (min)	m/z $[M+H]^+$
292	4-[6-(4-Fluorophenyl)-2-(3-hydroxypropyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridin-4-yl]phenol	Example 271	1	4.41	441.1
293	4-[6-(4-Fluorophenyl)-2-methyl-5-	Example 257	1	5.71	398.1

	pyrimidin-4-yl)pyrazolo[3,4- <i>b</i>]pyridin-4-yl]phenol				
294	3-[6-(4-Fluorophenyl)-2-(3-hydroxypropyl)-5-(4-pyridyl)pyrazolo[3,4- <i>b</i>]pyridin-4-yl]phenol	Example 277	1	4.54	441.2
295	3-[6-(4-Fluorophenyl)-2-methyl-5-(4-pyridyl)pyrazolo[3,4- <i>b</i>]pyridin-4-yl]phenol	Example 261	1	4.74	397.0

EXAMPLE 296**4,6-Bis(4-fluorophenyl)-5-(4-pyridyl)-2-(pyrrolidin-2-ylmethyl)pyrazolo[3,4-*b*]pyridine**

- 5 To a solution of 2-(1-benzylpyrrolidin-2-ylmethyl)-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine (83 mg, 0.1 mmol, obtained in example 259) in EtOH (6.9 mL), 10% Pd/C (8 mg) and formic acid (0.34 mL) were added and heated a reflux for 2 h. It was allowed to cool and filtered through celite, washed with EtOH and concentrated. The crude product obtained was purified by
- 10 chromatography on silica gel using hexane-EtOAc-MeOH-NH₃ of increasing polarity as eluent, to afford 40 mg of the title compound (yield: 57%)
- LC-MS (method 1): *t_R* = 4.98 min; *m/z* = 468.1 [M+H]⁺.

EXAMPLE 297**4-[4-[6-(4-Fluorophenyl)-2-methylpyrazolo[3,4-*b*]pyridin-5-yl]pyridin-2-ylamino]benzenesulfonamide**

- 15 Following a similar procedure to that described in example 214, but starting from 5-(2-chloropyridin-4-yl)-6-(4-fluorophenyl)-2-methylpyrazolo[3,4-*b*]pyridine (obtained in example 263) and 4-aminobenzenesulfonamide, the title compound was obtained.
- 20 LC-MS (method 1): *t_R* = 5.36 min; *m/z* = 475.3 [M+H]⁺.

EXAMPLE 298***N*-[5-[2-[(Cyclopropylmethyl)amino]pyrimidin-4-yl]-6-(4-fluorophenyl)-1*H*-pyrazolo[3,4-*b*]pyridin-3-yl]acetamide 7-oxide**

- Following a similar procedure to that described in example 208, but starting from
- 25 *N*-[6-(4-fluorophenyl)-5-(2-methylsulfonylpyrimidin-4-yl)-1*H*-pyrazolo[3,4-*b*]pyridin-

3-yl]acetamide 7-oxide (obtained in example 268 section a) and (cyclopropylmethyl)amine, the title compound was obtained.

LC-MS (method 1): $t_R = 5.45$ min; $m/z = 434.2$ $[M+H]^+$.

EXAMPLE 299

5 ***N*-[6-(4-Fluorophenyl)-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridin-3-yl]isonicotinamide**

a) Isonicotinoyl chloride hydrochloride

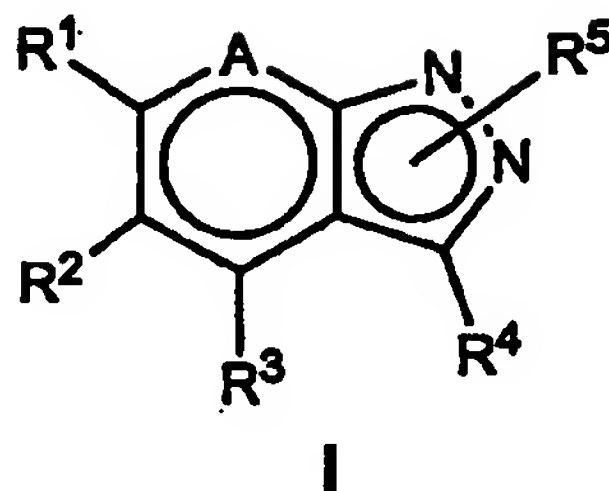
A solution of isonicotinic acid (0.10 g, 0.8 mmol) and thionyl chloride (1 mL) were heated to reflux for 2 h and concentrated. The product obtained was used
10 immediately in the following step.

b) Title compound

In a volumetric flask, 3-amino-6-(4-fluorophenyl)-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine (0.20 g, 0.7 mmol, obtained in example 70), isonicotinoyl chloride hydrochloride (0.12 g, 0.7 mmol, obtained in section a) and pyridine (1 mL) were
15 introduced under argon atmosphere. This was stirred at room temperature for 2 days. It was concentrated and the residue dissolved in a mixture of $CHCl_3$ and 1 N HCl. The phases were separated and the aqueous phase was extracted with $CHCl_3$ (x2). The aqueous phase was basified by slow addition of 1 N NaOH. Brine was added and extracted with $CHCl_3$ and EtOAc. The organic phase was dried
20 over Na_2SO_4 and concentrated to dryness. The crude product obtained was purified by chromatography on silica gel using $CHCl_3$ -MeOH mixtures of increasing polarity as eluent, to afford 98 mg of the title compound (yield: 68%).
LC-MS (method 1): $t_R = 4.21$ min; $m/z = 411.1$ $[M+H]^+$.

CLAIMS

1.- A compound of general formula I



5

wherein:

A represents N or N⁺O⁻;

10 R¹ represents phenyl or Het optionally substituted with one or more substituents selected from R^a and R^b;

R² represents Het optionally substituted with one or more substituents selected from R^a and R^b;

15 R³ represents H, Cy optionally substituted with one or more substituents selected from R^a and R^b, or R³ represents C₁₋₆alkyl optionally substituted with one or more substituents selected from R^b and Cy*, wherein Cy* can be optionally substituted with one or more substituents selected from R^b and R^c;

20 R⁴ represents H, R^a, halogen, -OR^{a'}, -OCOR^a, -OSO₂R^a, -OCONR^aR^{a'}, -NO₂, -CN, -COR^{a'}, -CO₂R^a, -CONR^{a'}R^{a'}, -NR^{a'}R^{a'}, -NR^{a'}COR^{a'}, -NR^{a'}CONR^{a'}R^{a'}, -NR^{a'}CO₂R^a, -NR^{a'}SO₂R^a, -SR^{a'}, -SOR^a, -SO₂R^a or -SO₂NR^{a'}R^{a'};

25 R⁵ can be placed on any of the 2 N of the pyrazole ring of formula I and represents H or R^f;

each R^a independently represents C₁₋₆alkyl, C₂₋₆alkenyl, C₂₋₆alkynyl or Cy, wherein the groups C₁₋₆alkyl, C₂₋₆alkenyl or C₂₋₆alkynyl can be optionally substituted with

one or more substituents selected from R^b and Cy^* , and wherein any of the groups Cy or Cy^* can be optionally substituted with one or more substituents selected from R^b and R^c ;

5 each R^a independently represents H or R^a ;

each R^b independently represents halogen, $-OR^c$, $-OCOR^c$, $-OSO_2R^c$, $-OCONR^cR^c$, $-NO_2$, $-CN$, $-COR^c$, $-CO_2R^c$, $-CONR^cR^c$, $-CONR^cNR^cR^c$, $-NR^cR^c$, $-NR^cCOR^c$, $-NR^cCONR^cR^c$, $-NR^cCO_2R^c$, $-NR^cSO_2R^c$, $-SR^c$, $-SOR^c$,
 10 $-SO_2R^c$, $-SO_2NR^cR^c$, $-C(NR^c)NR^cR^c$, $-C(NSO_2NR^cR^c)NR^cR^c$, $-C(NOR^c)R^c$, $-C(NNR^cR^c)R^c$, $-NR^cC(NR^c)NR^cR^c$ or $-NR^cC(NCN)NR^cR^c$;

each R^c independently represents C_{1-6} alkyl, C_{2-6} alkenyl, C_{2-6} alkynyl or Cy , wherein all these groups can be optionally substituted with one or more substituents R^d ;

15

each R^c independently represents H or R^c ;

each R^d independently represents halogen, R^e , $-OR^e$, $-OCOR^e$, $-OSO_2R^e$, $-OCONR^eR^e$, $-NO_2$, $-CN$, $-COR^e$, $-CO_2R^e$, $-CONR^eR^e$, $-CONR^eNR^eR^e$,
 20 $-NR^eR^e$, $-NR^eCOR^e$, $-NR^eCONR^eR^e$, $-NR^eCO_2R^e$, $-NR^eSO_2R^e$, $-SR^e$, $-SOR^e$, $-SO_2R^e$, $-SO_2NR^eR^e$, $-C(NR^e)NR^eR^e$, $-C(NSO_2NR^eR^e)NR^eR^e$, $-C(NOR^e)R^e$, $-C(NNR^eR^e)R^e$, $-NR^eC(NR^e)NR^eR^e$, $-NR^eC(NCN)NR^eR^e$ or Cy optionally substituted with one or more substituents selected from halogen, R^e , $-OR^e$, $-OCOR^e$, $-OSO_2R^e$, $-OCONR^eR^e$, $-NO_2$, $-CN$, $-COR^e$, $-CO_2R^e$, $-CONR^eR^e$,
 25 $-CONR^eNR^eR^e$, $-NR^eR^e$, $-NR^eCOR^e$, $-NR^eCONR^eR^e$, $-NR^eCO_2R^e$, $-NR^eSO_2R^e$, $-SR^e$, $-SOR^e$, $-SO_2R^e$, $-SO_2NR^eR^e$, $-C(NR^e)NR^eR^e$, $-C(NSO_2NR^eR^e)NR^eR^e$, $-C(NOR^e)R^e$, $-C(NNR^eR^e)R^e$, $-NR^eC(NR^e)NR^eR^e$ and $-NR^eC(NCN)NR^eR^e$;

30 each R^e independently represents C_{1-6} alkyl or halo C_{1-6} alkyl;

each R^e independently represents H or R^e ;

R^f represents C_{1-6} alkyl, C_{2-6} alkenyl, C_{2-6} alkynyl or Cy, wherein the groups C_{1-6} alkyl, C_{2-6} alkenyl or C_{2-6} alkynyl can be optionally substituted with one or more substituents selected from R^g and Cy^* , and wherein any of the groups Cy or Cy^* can be optionally substituted with one or more substituents selected from R^g and R^a ;

each R^g independently represents halogen, $-OR^a$, $-OCOR^a$, $-OSO_2R^a$, $-OCONR^aR^a$, $-NO_2$, $-CN$, $-COR^a$, $-CO_2R^a$, $-CONR^aR^a$, $-CONR^aNR^aR^a$, $-NR^aR^a$, $-NR^aCOR^a$, $-NR^aCONR^aR^a$, $-NR^aCO_2R^a$, $-NR^aSO_2R^a$, $-SR^a$, $-SOR^a$, $-SO_2R^a$, $-SO_2NR^aR^a$, $-C(NR^a)NR^aR^a$, $-C(NSO_2NR^aR^a)NR^aR^a$, $-C(NOR^a)R^a$, $-C(NNR^aR^a)R^a$, $-NR^aC(NR^a)NR^aR^a$ or $-NR^aC(NCN)NR^aR^a$;

Het in the above definitions represents pyridine, pyrazine, pyrimidine, pyridazine, 2(1H)-pyridone, 2(1H)-pyrazinone, 2(1H)-pyrimidinone or 2(1H)-pyridazinone;

15

Cy or Cy^* in the above definitions represent a partially unsaturated, saturated or aromatic 3- to 7-membered monocyclic or 8- to 12-membered bicyclic carbocyclic ring, which optionally contains from 1 to 4 heteroatoms selected from N, S and O, which can optionally contain 1 or 2 oxo groups when the ring is saturated or partially unsaturated, and wherein said ring or rings can be bonded to the rest of the molecule through a carbon or a nitrogen atom; and the salts, solvates and prodrugs thereof.

20

2.- A compound according to claim 1 wherein A represents N.

3.- A compound according to claim 2 wherein R^b can be placed on any of the 2 N of the pyrazole ring of formula I and represents H or R^a .

25

4.- A compound according to any of claims 1 to 3 with the proviso that when R^3 and R^5 both represent H and R^2 represents Het optionally substituted with one or more substituents selected from halogen, $-CN$, $-CF_3$, $-OH$, $-NO_2$, $-OR^b$, $-NR^bR^b$, $-OCF_3$, C_{1-6} alkyl, C_{2-6} alkenyl, C_{2-6} alkynyl and Cy, wherein Cy can be optionally substituted with one or more substituents selected from R^b and R^c , and wherein R^b represents C_{1-6} alkyl, then R^4 is not $-NR^aCOR^a$, $-NHCONHR^a$ or $-NHCO_2R^a$.

30

- 5.- A compound according to any of claims 1 to 4 wherein R^1 represents pyridine or phenyl, wherein all these groups can be optionally substituted with one or more substituents selected from R^a and R^b .
- 6.- A compound according to claim 5 wherein R^1 represents phenyl optionally substituted with one or more substituents selected from R^a and R^b .
- 7.- A compound according to claim 6 wherein R^1 represents phenyl optionally substituted with one or more substituents selected from halogen, $-OR^c$, $-NO_2$, $-CN$, $-CONR^cR^c$, $-NR^cR^c$ and C_{1-6} alkyl optionally substituted with one or more substituents selected from halogen, $-OR^c$, $-COR^c$, $-NR^cR^c$ and $-NR^cCOR^c$.
- 8.- A compound according to claim 7 wherein R^1 represents phenyl optionally substituted with one or more substituents selected from halogen and halo C_{1-6} alkyl.
- 9.- A compound according to any of claims 1 to 8 wherein R^2 represents pyridine or pyrimidine, wherein all these groups can be optionally substituted with one or more substituents selected from R^a and R^b .
- 10.- A compound according to claim 9 wherein R^2 represents 4-pyridine or 4-pyrimidine, wherein all these groups can be optionally substituted with one or more substituents selected from R^a and R^b .
- 11.- A compound according to claim 10 wherein R^2 represents 4-pyridine or 4-pyrimidine, wherein all these groups can be optionally substituted with one or more substituents selected from halogen, $-OR^c$, $-NR^cR^c$, $-SR^c$ and $-SO_2R^c$.
- 12.- A compound according to claim 11 wherein R^2 represents 4-pyridine.
- 13.- A compound according to claim 11 wherein R^2 represents 4-pyrimidine substituted with $-NR^cR^c$, wherein in R^2 :
each R^c independently represents H or R^c ;
each R^c independently represents C_{1-6} alkyl optionally substituted with one or more substituents selected from Cy and $-OR^e$; and
each R^e independently represents H or R^e .
- 14.- A compound according to any of claims 1 to 13 wherein R^3 represents H or Cy optionally substituted with one or more substituents selected from R^a and R^b .
- 15.- A compound according to claim 14 wherein R^3 represents H, heteroaryl or phenyl, wherein all these groups can be optionally substituted with one or more substituents selected from R^a and R^b , and wherein heteroaryl represents an aromatic 5- or 6-membered monocyclic or 8- to 12-membered bicyclic ring, which

contains from 1 to 4 heteroatoms selected from N, S and O and which can be bonded to the rest of the molecule through a carbon or a nitrogen atom.

16.- A compound according to claim 15 wherein R^3 represents heteroaryl or phenyl, wherein all these groups can be optionally substituted with one or more substituents selected from R^a and R^b .

17.- A compound according to claim 16 wherein R^3 represents monocyclic heteroaryl or phenyl, wherein all these groups can be optionally substituted with one or more substituents selected from halogen, $-NO_2$, $-OR^c$, C_{1-6} alkyl and Cy, wherein C_{1-6} alkyl can be optionally substituted with one or more substituents selected from R^b and Cy^* , and wherein any of the groups Cy or Cy^* can be optionally substituted with one or more substituents selected from R^b and R^c .

18.- A compound according to claim 17 wherein R^3 represents monocyclic heteroaryl or phenyl, wherein all these groups can be optionally substituted with one or more substituents selected from halogen, $-NO_2$, $-OR^c$, C_{1-6} alkyl, halo C_{1-6} alkyl and Cy, and wherein in R^3 :

each R^c independently represents H or R^c ;

each R^c independently represents C_{1-6} alkyl optionally substituted with one or more substituents R^d ; and

each R^d independently represents Cy.

19.- A compound according to any of claims 1 to 18 wherein R^4 represents H, R^a , halogen, $-OR^a$, $-CN$, $-CONR^aR^a$, $-NR^aR^a$ or $-NR^aCOR^a$.

20.- A compound according to claim 19 wherein R^4 represents H.

21.- A compound according to any of claims 1 to 20 wherein R^5 represents H or R^5 represents R^a and is placed on the N at position 2 of the pyrazole ring.

22.- A compound according to claim 21 wherein R^5 represents R^a .

23.- A compound according to claim 21 or 22 wherein R^b in R^5 represents C_{1-6} alkyl, C_{2-6} alkenyl or Cy, wherein the groups C_{1-6} alkyl or C_{2-6} alkenyl can be optionally substituted with one or more substituents selected from R^b and Cy^* , and wherein any of the groups Cy or Cy^* can be optionally substituted with one or more substituents selected from R^b and R^c .

24.- A compound according to claim 23 wherein each R^b in R^5 independently represents halogen, $-OR^c$, $-COR^c$, $-CO_2R^c$, $-CONR^cR^c$, $-NR^cR^c$, $-NR^cCOR^c$, $-NR^cCONR^cR^c$, $-NR^cSO_2R^c$, $-SR^c$, $-SOR^c$ or $-SO_2R^c$.

25.- A compound according to claim 24 wherein in R⁵:

each R^b independently represents -OR^c, -COR^c, -CONR^cR^c, -NR^cR^c, -NR^cCOR^c, -NR^cCONR^cR^c, -NR^cSO₂R^c, -SOR^c or -SO₂R^c;

each R^c independently represents H or R^e;

5 each R^o independently represents Cy or C₁₋₆alkyl, wherein all these groups can be optionally substituted with one or more substituents selected from R^d;

each R^d independently represents R^o, -OR^o, -NR^oR^o, -CN, -COR^o, -SR^o, -SOR^o or Cy.

10 26.- A compound according to claim 22 wherein R^a in R⁵ represents C₁₋₆alkyl optionally substituted with one or more substituents selected from -OR^c, -COR^c, -CONR^cR^c, -NR^cR^c, -NR^cCOR^c, -NR^cCONR^cR^c, -NR^cSO₂R^c and Cy* optionally substituted with one or more substituents selected from R^o; wherein in R⁵:

each R^c independently represents H or R^e;

15 each R^o independently represents Cy or C₁₋₆alkyl, wherein all these groups can be optionally substituted with one or more substituents selected from R^d;

each R^d independently represents -OR^o, -NR^oR^o, -CN, -COR^o, -SR^o, -SOR^o or Cy;

each R^e independently represents H or R^e; and

each R^o independently represents C₁₋₆alkyl.

20 27.- A compound according to claim 1 selected from:

4,6-bis(4-fluorophenyl)-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine;

4,6-diphenyl-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine;

5-(4-pyridyl)-4,6-bis[3-(trifluoromethyl)phenyl]-1*H*-pyrazolo[3,4-*b*]pyridine;

4,6-bis(4-fluorophenyl)-3-methyl-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine;

25 3-methyl-4,6-diphenyl-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine;

2-ethyl-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;

1-ethyl-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;

4,6-bis(4-fluorophenyl)-2-methyl-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;

4,6-bis(4-fluorophenyl)-1-methyl-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;

30 4,6-bis(4-fluorophenyl)-2,3-dimethyl-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;

4,6-bis(4-fluorophenyl)-1,3-dimethyl-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;

2-[2-[1-(*tert*-butoxycarbonyl)piperidin-4-yl]ethyl]-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;

- 1-[2-[1-(*tert*-butoxycarbonyl)piperidin-4-yl]ethyl]-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;
2-[1-(*tert*-butoxycarbonyl)piperidin-4-yl]-4,6-bis(4-fluorophenyl)-3-methyl-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;
5 1-[1-(*tert*-butoxycarbonyl)piperidin-4-yl]-4,6-bis(4-fluorophenyl)-3-methyl-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;
2-(3-chloropropyl)-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;
1-(3-chloropropyl)-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;
3-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]propan-1-ol;
10 3-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-1-yl]propan-1-ol;
2-[1-(*tert*-butoxycarbonyl)piperidin-4-yl]-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;
1-[1-(*tert*-butoxycarbonyl)piperidin-4-yl]-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;
15 2-methyl-4,6-diphenyl-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;
1-methyl-4,6-diphenyl-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;
4,6-bis(4-fluorophenyl)-5-(4-pyridyl)-2-[2-(tetrahydropyran-2-yloxy)ethyl]-pyrazolo[3,4-*b*]pyridine;
2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-1-yl]ethanol;
20 4,6-bis(4-fluorophenyl)-2-(4-methylsulfanylbenzyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;
4,6-bis(4-fluorophenyl)-1-(4-methylsulfanylbenzyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;
2-[1-(*tert*-butoxycarbonyl)piperidin-4-ylmethyl]-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;
25 1-[1-(*tert*-butoxycarbonyl)piperidin-4-ylmethyl]-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;
4,6-bis(4-fluorophenyl)-2-[2-(morpholin-4-yl)ethyl]-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;
30 4,6-bis(4-fluorophenyl)-1-[2-(morpholin-4-yl)ethyl]-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;
ethyl 2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]acetate;
ethyl 2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-1-yl]acetate;

- ethyl 3-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]propionate;
ethyl 3-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-1-yl]propionate;
4,6-bis(4-fluorophenyl)-2-(4-piperidyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;
4,6-bis(4-fluorophenyl)-1-(4-piperidyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;
5 4,6-bis(4-fluorophenyl)-2-(4-piperidylmethyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;
4,6-bis(4-fluorophenyl)-1-(4-piperidylmethyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;
4,6-bis(6-chloropyridin-3-yl)-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine;
4,6-bis(4-fluorophenyl)-3-methyl-2-(4-piperidyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;
10 4,6-bis(4-fluorophenyl)-3-methyl-1-(4-piperidyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;
4,6-bis(4-fluorophenyl)-2-[2-(4-piperidyl)ethyl]-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;
2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]acetic acid;
2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-1-yl]acetic acid;
15 3-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]proplonic acid;
3-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-1-yl]propionic acid;
2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]-1-(morpholin-4-yl)ethanone;
2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-1-yl]acetamide;
20 2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-1-yl]-1-(morpholin-4-yl)ethanone;
3-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]-1-(morpholin-4-yl)propan-1-one;
3-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-1-yl]-*N*-
25 propylpropionamide;
3-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-1-yl]-1-(morpholin-4-yl)propan-1-one;
4,6-bis(4-fluorophenyl)-2-(4-methylsulfanylphenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;
30 4,6-bis(4-fluorophenyl)-1-(4-methylsulfanylphenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;
4,6-bis(4-fluorophenyl)-2-(4-methylsulfinylphenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;

- 4,6-bis(4-fluorophenyl)-2-(4-methylsulfonylphenyl)-5-(4-pyridyl)pyrazolo[3,4-
b]pyridine;
4,6-bis(4-fluorophenyl)-1-(4-methylsulfinylphenyl)-5-(4-pyridyl)pyrazolo[3,4-
b]pyridine;
5 4,6-bis(4-fluorophenyl)-2-(4-methylsulfinylbenzyl)-5-(4-pyridyl)pyrazolo[3,4-
b]pyridine;
4,6-bis(4-fluorophenyl)-2-(4-methylsulfonylbenzyl)-5-(4-pyridyl)pyrazolo[3,4-
b]pyridine;
4,6-bis(4-fluorophenyl)-1-(4-methylsulfinylbenzyl)-5-(4-pyridyl)pyrazolo[3,4-
10 b]pyridine;
4,6-bis(4-fluorophenyl)-1-(4-methylsulfonylbenzyl)-5-(4-pyridyl)pyrazolo[3,4-
b]pyridine;
3-chloro-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)-1H-pyrazolo[3,4-b]pyridine;
3-bromo-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)-1H-pyrazolo[3,4-b]pyridine;
15 4,6-bis(4-fluorophenyl)-5-(4-pyridyl)-1H-pyrazolo[3,4-b]pyridine-3-carbonitrile;
3-bromo-4,6-bis(4-fluorophenyl)-1-methyl-5-(4-pyridyl)pyrazolo[3,4-b]pyridine;
4,6-bis(4-fluorophenyl)-5-(4-pyridyl)-1H-pyrazolo[3,4-b]pyridine-3-carboxamide;
3-aminomethyl-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)-1H-pyrazolo[3,4-b]pyridine;
4,6-bis(4-fluoro-3-nitrophenyl)-5-(4-pyridyl)-1H-pyrazolo[3,4-b]pyridine;
20 3-amino-6-(4-fluorophenyl)-5-(4-pyridyl)-1H-pyrazolo[3,4-b]pyridine;
3-amino-6-(4-fluorophenyl)-1-methyl-5-(4-pyridyl)pyrazolo[3,4-b]pyridine;
4-[6-(4-fluorophenyl)-5-(4-pyridyl)-1H-pyrazolo[3,4-b]pyridin-4-yl]phenol;
2-(2,2-diethoxyethyl)-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridine;
1-(2,2-diethoxyethyl)-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridine;
25 4,6-bis(4-fluorophenyl)-1-methyl-5-(4-pyridyl)pyrazolo[3,4-b]pyridine-3-carbonitrile;
3-bromo-6-(4-fluorophenyl)-5-(4-pyridyl)-1H-pyrazolo[3,4-b]pyridine;
6-fluorophenyl-5-(4-pyridyl)-1H-pyrazolo[3,4-b]pyridine;
N-methyl-[3-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)-pyrazolo[3,4-b]pyridin-2-
yl]propyl]amine;
30 [4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridin-2-yl]methanol;
2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-b]pyridin-2-yl]-N,N-
dimethylacetamide;

- 2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-1-yl]-*N,N*-dimethylacetamide;
- 4,6-bis(4-fluorophenyl)-2-[2-(2-methoxyethoxy)ethyl]-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;
- 5 4,6-bis(4-fluorophenyl)-1-[2-(2-methoxyethoxy)ethyl]-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;
- 4,6-bis(4-fluorophenyl)-2-[3-(morpholin-4-yl)propyl]-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;
- 4,6-bis(6-chloropyridin-3-yl)-2-methyl-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;
- 10 4,6-bis(6-chloropyridin-3-yl)-1-methyl-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;
- 4,6-bis(6-chloropyridin-3-yl)-3-methyl-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine;
- 4,6-bis(6-methylpyridin-3-yl)-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine;
- 4,6-bis(4-fluorophenyl)-2-(2-phthalimidoethyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;
- 2-(2-aminoethyl)-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;
- 15 2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethanol;
- 6-(4-fluorophenyl)-2-methyl-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;
- 4,6-bis(4-fluorophenyl)-2-(3-phthalimidopropyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;
- 2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]acetaldehyde;
- 2-(3-aminopropyl)-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;
- 20 *N*-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridin-3-ylmethyl]-1-(*tert*-butoxycarbonyl)piperidine-4-carboxamide;
- N*-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridin-3-ylmethyl]-1*H*-piperidine-4-carboxamide;
- 2-(3-benzyloxypropyl)-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;
- 25 1-(3-benzyloxypropyl)-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;
- N,N*-diethyl-[2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl]amine;
- N,N*-diethyl-[2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-1-yl]ethyl]amine;
- 30 4,6-bis(4-fluorophenyl)-5-(4-pyridyl)-2-(3-pyridylmethyl)pyrazolo[3,4-*b*]pyridine;
- 4,6-bis(4-fluorophenyl)-5-(4-pyridyl)-1-pyridin-3-ylmethylpyrazolo[3,4-*b*]pyridine;
- N,N*-dimethyl-[3-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)-pyrazolo[3,4-*b*]pyridin-2-yl]propyl]amine;

- N,N*-dimethyl-[3-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-1-yl]propyl]amine;
1-[2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl]piperidin-4-ol;
5 3-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]-2-hydroxypropan-1-ol;
3-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-1-yl]-2-hydroxypropan-1-ol;
4,6-bis(4-fluorophenyl)-5-(4-pyridyl)-2-(4-pyridylmethyl)pyrazolo[3,4-*b*]pyridine;
10 4,6-bis(4-fluorophenyl)-5-(4-pyridyl)-1-(4-pyridylmethyl)pyrazolo[3,4-*b*]pyridine;
N-(*tert*-butoxycarbonyl)-[1-[3-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]propyl]piperidin-4-yl]amine;
2-[1-(*tert*-butoxycarbonyl)piperidin-4-yl]-6-(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;
15 1-[1-(*tert*-butoxycarbonyl)piperidin-4-yl]-6-(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;
3-methyl-4,6-bis(6-methylpyridin-3-yl)-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine;
1-[3-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]-propyl]piperidin-4-one;
20 *N*-(*tert*-butoxycarbonyl)-[1-[2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl]piperidin-4-yl]amine;
N-methyl-[1-[2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl]piperidin-4-yl]amine;
[1-[3-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]propyl]piperidin-4-yl]amine;
25 2-[1-[2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl]piperidin-4-yl]ethanol;
[1-[2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl]piperidin-4-yl]amine;
30 6-(4-fluorophenyl)-2-(4-piperidyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;
6-(4-fluorophenyl)-1-(4-piperidyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;
3-amino-5-[2-(methylsulfanyl)pyrimidin-4-yl]-6-[3-(trifluoromethyl)phenyl]-1*H*-pyrazolo[3,4-*b*]pyridine;

- 2-[3-[1-(4-*tert*-butoxycarbonyl)piperazin-1-yl]propyl]-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;
- 4,6-bis(4-fluorophenyl)-2-[3-(1-piperazin-1-yl)propyl]-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;
- 5 5-[2-(methylsulfanyl)pyrimidin-4-yl]-6-[3-(trifluoromethyl)phenyl]-1*H*-pyrazolo[3,4-*b*]pyridine;
- 5-[2-(methylsulfonyl)pyrimidin-4-yl]-6-[3-(trifluoromethyl)phenyl]-1*H*-pyrazolo[3,4-*b*]pyridine;
- (1*S*)-*N*-(1-phenylethyl)-[4-[6-[3-(trifluoromethyl)phenyl]-1*H*-pyrazolo[3,4-*b*]pyridin-5-yl]pyrimidin-2-yl]amine;
- 10 1-[3-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]propyl]piperidin-4-ol;
- 2-[1-[3-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]propyl]piperidin-4-yl]ethanol;
- 15 4,6-bis(4-fluorophenyl)-3-(4-piperidyl)-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine;
- 6-(4-fluorophenyl)-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine-3-carbonitrile;
- 2-[2-[[1-(*tert*-butoxycarbonyl)piperidin-4-yl]amino]ethyl]-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;
- 4,6-bis(4-fluorophenyl)-2-[2-[(4-piperidyl)amino]ethyl]-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;
- 20 *N*-(2-methoxyethyl)-[2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl]amine;
- 1-[4-[2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl]piperazin-1-yl]ethanone;
- 25 3-[4,6-diphenyl-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]propan-1-ol;
- 3-[4,6-diphenyl-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-1-yl]propan-1-ol;
- 2-ethyl-4,6-diphenyl-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;
- 1-ethyl-4,6-diphenyl-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;
- 4,6-diphenyl-2-(2-phthalimidoethyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;
- 30 2-(2-aminoethyl)-4,6-diphenyl-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;
- 2-allyl-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;
- 1-allyl-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;

- 1-[2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl]piperidin-4-one;
- 3-aminomethyl-6-(4-fluorophenyl)-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine;
- 3-amino-6-(4-fluorophenyl)-4-methyl-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine;
- 5 3-[*N*-[2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl]amino]propan-1-ol;
- N*-ethyl-[2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl]amine;
- 2-[*N*-[2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl]amino]ethanol;
- 10 *N*-[(2-pyridyl)methyl]-[2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl]amine;
- N*-[(2-thienyl)methyl]-[2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl]amine;
- 15 1-[2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl]piperidine-4-carboxamide;
- 4,6-bis(4-fluorophenyl)-2-[2-(pyrrolidin-1-yl)ethyl]-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;
- (3*R*)-1-[2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl]pyrrolidin-3-ol;
- 20 2-[*N*-[2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl]-*N*-methylamino]ethanol;
- 4,6-bis(4-fluorophenyl)-5-(4-pyridyl)-2-[2-(1,2,3,4-tetrahydroisoquinolin-2-yl)ethyl]pyrazolo[3,4-*b*]pyridine;
- 25 4,6-bis(4-fluorophenyl)-2-[2-(4-phenylpiperazin-1-yl)ethyl]-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;
- 4,6-bis(4-fluorophenyl)-2-[2-[4-(1-piperidyl)piperidin-1-yl]ethyl]-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;
- 3-[*N*-[2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl]-*N*-methylamino]propiononitrile;
- 30 *N*-methyl-[2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl]amine;

- 2-[2-[4-(*tert*-butoxycarbonyl)piperazin-1-yl]ethyl]-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;
4,6-bis(4-fluorophenyl)-2-[2-(piperazin-1-yl)ethyl]-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;
5 4,6-bis(4-fluorophenyl)-5-(4-pyridyl)-2-vinylpyrazolo[3,4-*b*]pyridine;
2-[*N*-{2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl}-*N*-(2-hydroxyethyl)amino]ethanol;
N-cyclopropyl-[2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl]amine;
10 *N*-[2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl]acetamide;
N-[2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl]-*n*'-isopropylurea;
N-[2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl]methanesulfonamide;
15 6-(4-fluorophenyl)-4-(4-piperidyl)-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine;
6-(4-fluorophenyl)-4-(2-furyl)-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine;
6-(4-fluorophenyl)-4-(1*H*-imidazol-4-yl)-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine;
4-(5-bromothien-2-yl)-6-(4-fluorophenyl)-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine;
20 4,6-bis(4-fluorophenyl)-5-(2-methylsulfanylpirimidin-4-yl)-1*H*-pyrazolo[3,4-*b*]pyridine;
5-(2-chloropyridin-4-yl)-4,6-bis(4-fluorophenyl)-1*H*-pyrazolo[3,4-*b*]pyridine;
6-(4-fluorophenyl)-4-(2-phenylethyl)-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine;
4-(6-chloropyridin-3-yl)-6-(4-fluorophenyl)-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine;
25 4-(3,4-dichlorophenyl)-1-ethyl-6-(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;
6-(4-fluorophenyl)-4-(1-methylpiperidin-4-yl)-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine;
3-amino-6-(4-fluorophenyl)-5-(2-methylsulfanylpirimidin-4-yl)-1*H*-pyrazolo[3,4-*b*]pyridine;
30 6-(4-fluorophenyl)-5-(2-methylsulfanylpirimidin-4-yl)-1*H*-pyrazolo[3,4-*b*]pyridine;
4,6-diphenyl-5-(4-pyridyl)-2-[2-(tetrahydropyran-2-yloxy)ethyl]pyrazolo[3,4-*b*]pyridine;

- 6-(4-fluorophenyl)-4-(2-furyl)-2-methyl-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;
6-(4-fluorophenyl)-2-methyl-4-(1-methyl-1*H*-imidazol-4-yl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;
6-(4-fluorophenyl)-5-(2-methylsulfanylpirimidin-4-yl)-2-[2-(tetrahydropyran-2-yloxy)ethyl]pyrazolo[3,4-*b*]pyridine;
5 6-(4-fluorophenyl)-5-(2-methylsulfanylpirimidin-4-yl)-2-[3-(tetrahydropyran-2-yloxy)propyl]pyrazolo[3,4-*b*]pyridine;
4,6-bis(4-fluorophenyl)-5-(2-methylsulfanylpirimidin-4-yl)-2-[3-(tetrahydropyran-2-yloxy)propyl]pyrazolo[3,4-*b*]pyridine;
10 4-(5-bromothien-2-yl)-6-(4-fluorophenyl)-2-methyl-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine;
6-(4-fluorophenyl)-2-methyl-5-(2-methylsulfanylpirimidin-4-yl)pyrazolo[3,4-*b*]pyridine;
5-(2-chloropyridin-4-yl)-4,6-bis(4-fluorophenyl)-2-methylpyrazolo[3,4-*b*]pyridine;
15 6-(4-fluorophenyl)-4-(2-phenylethyl)-5-(4-pyridyl)-2-[3-(tetrahydropyran-2-yloxy)propyl]pyrazolo[3,4-*b*]pyridine;
4-(6-chloropyridin-3-yl)-6-(4-fluorophenyl)-5-(4-pyridyl)-2-[3-(tetrahydropyran-2-yloxy)propyl]pyrazolo[3,4-*b*]pyridine;
4-(6-chloropyridin-3-yl)-6-(4-fluorophenyl)-2-methyl-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;
20 5-(2-methylsulfanylpirimidin-4-yl)-2-[3-(tetrahydropyran-2-yloxy)propyl]-6-(3-trifluoromethylphenyl)pyrazolo[3,4-*b*]pyridine;
6-(4-fluorophenyl)-2-methyl-5-(4-pyridyl)-4-[5-(3-pyridyl)thien-2-yl]pyrazolo[3,4-*b*]pyridine;
25 2-[4,6-diphenyl-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethanol;
3-[5-(2-methylsulfanylpirimidin-4-yl)-6-(3-trifluoromethylphenyl)pyrazolo[3,4-*b*]pyridin-2-yl]propan-1-ol;
2-[6-(4-fluorophenyl)-5-(2-methylsulfanylpirimidin-4-yl)pyrazolo[3,4-*b*]pyridin-2-yl]ethanol;
30 3-[6-(4-fluorophenyl)-5-(2-methylsulfanylpirimidin-4-yl)pyrazolo[3,4-*b*]pyridin-2-yl]propan-1-ol;
3-[4,6-bis(4-fluorophenyl)-5-(2-methylsulfanylpirimidin-4-yl)pyrazolo[3,4-*b*]pyridin-2-yl]propan-1-ol;

- 3-[6-(4-fluorophenyl)-4-(2-phenylethyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]propan-1-ol;
3-[4-(6-chloropyridin-3-yl)-6-(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]propan-1-ol;
5 6-(4-fluorophenyl)-2-methyl-5-(2-methylsulfonylpyrimidin-4-yl)pyrazolo[3,4-*b*]pyridine;
3-[5-(2-methylsulfonylpyrimidin-4-yl)-6-(3-trifluoromethylphenyl)pyrazolo[3,4-*b*]pyridin-2-yl]propan-1-ol;
2-[6-(4-fluorophenyl)-5-(2-methylsulfonylpyrimidin-4-yl)pyrazolo[3,4-*b*]pyridin-2-yl]ethanol;
10 3-[6-(4-fluorophenyl)-5-(2-methylsulfonylpyrimidin-4-yl)pyrazolo[3,4-*b*]pyridin-2-yl]propan-1-ol;
3-[4,6-bis(4-fluorophenyl)-5-(2-methylsulfonylpyrimidin-4-yl)pyrazolo[3,4-*b*]pyridin-2-yl]propan-1-ol;
15 *N*-cyclopropylmethyl-[4-[6-[3-(trifluoromethyl)phenyl]-1*H*-pyrazolo[3,4-*b*]pyridin-5-yl]pyrimidin-2-yl]amine;
(1*S*)-3-[5-[2-(1-phenylethylamino)pyrimidin-4-yl]-6-(3-trifluoromethylphenyl)pyrazolo[3,4-*b*]pyridin-2-yl]propan-1-ol;
N-cyclopropylmethyl-[4-[6-(4-fluorophenyl)-2-methylpyrazolo[3,4-*b*]pyridin-5-yl]pyrimidin-2-yl]amine;
20 2-[5-[2-[(cyclopropylmethyl)amino]pyrimidin-4-yl]-6-(4-fluorophenyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethanol;
3-[5-[2-[(cyclopropylmethyl)amino]pyrimidin-4-yl]-6-(4-fluorophenyl)pyrazolo[3,4-*b*]pyridin-2-yl]propan-1-ol;
25 3-[5-[2-[(cyclopropylmethyl)amino]pyrimidin-4-yl]-4,6-bis(4-fluorophenyl)pyrazolo[3,4-*b*]pyridin-2-yl]propan-1-ol;
4-[4-[4,6-bis(4-fluorophenyl)-2-methylpyrazolo[3,4-*b*]pyridin-5-yl]pyridin-2-ylamino]benzenesulfonamide;
4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-3-ol;
30 6-(4-fluorophenyl)-4-(3*H*-imidazol-4-yl)-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine;
6-(4-fluorophenyl)-4-(1*H*-pyrazol-3-yl)-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine;
3-[6-(4-fluorophenyl)-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridin-4-yl]phenol;
4-cyclopropyl-6-(4-fluorophenyl)-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine;

- 6-(4-fluorophenyl)-4-(5-methylfuran-2-yl)-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine;
4-(5-bromofuran-2-yl)-6-(4-fluorophenyl)-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine;
4-(4-benzyloxyphenyl)-6-(4-fluorophenyl)-5-pyrimidin-4-yl-1*H*-pyrazolo[3,4-*b*]pyridine;
5 4-(4-benzyloxyphenyl)-6-(4-fluorophenyl)-5-(2-methylsulfanylpymidin-4-yl)-1*H*-pyrazolo[3,4-*b*]pyridine;
6-(4-fluorophenyl)-4-propyl-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine;
4-(3-benzyloxyphenyl)-6-(4-fluorophenyl)-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine;
5-(2-chloropyridin-4-yl)-6-(4-fluorophenyl)-1*H*-pyrazolo[3,4-*b*]pyridine;
10 4-[6-(4-fluorophenyl)-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridin-4-yl]butan-1-ol;
4-benzyl-6-(4-fluorophenyl)-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine;
4-(4-benzyloxyphenyl)-6-(4-fluorophenyl)-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridine;
4,6-bis(4-fluorophenyl)-5-pyrimidin-4-yl-1*H*-pyrazolo[3,4-*b*]pyridine;
[(2*S*)-2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl]pyrrolidine-2-carboxamide;
15 2-[2-(4,6-diphenyl-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl)ethylamino]ethanol;
6-(4-fluorophenyl)-2-methyl-4-(3-pyridyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;
6-(4-fluorophenyl)-3-methyl-5-(2-methylsulfanylpymidin-4-yl)-1*H*-pyrazolo[3,4-*b*]pyridine;
20 3-amino-5-(2-methylsulfanylpymidin-4-yl)-6-phenyl-1*H*-pyrazolo[3,4-*b*]pyridine;
5-(2-methylsulfanylpymidin-4-yl)-6-phenyl-1*H*-pyrazolo[3,4-*b*]pyridine;
6-(4-fluorophenyl)-4-methyl-5-(2-methylsulfanylpymidin-4-yl)-1*H*-pyrazolo[3,4-*b*]pyridine;
5-(2-methoxypymidin-4-yl)-6-(3-trifluoromethylphenyl)-1*H*-pyrazolo[3,4-*b*]pyridine;
25 *N*-[2-[4,6-(di)phenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl]acetamide;
N-[3-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]propyl]acetamide;
N-[2-(4,6-diphenyl-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl)ethyl]-*N*-(2-hydroxyethyl)acetamide;
30 *N*-[2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl]propionamide;
N-[3-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]propyl]methanesulfonamide;

- 5-(2-aminopyrimidin-4-yl)-6-(3-trifluoromethylphenyl)-1*H*-pyrazolo[3,4-*b*]pyridine;
N-[5-(2-methylsulfanylpymidin-4-yl)-6-(3-trifluoromethylphenyl)-1*H*-pyrazolo[3,4-*b*]pyridin-3-yl]acetamide;
N-cyclopropylmethyl-[4-[3-benzyloxycarbonylamino-6-(3-trifluoromethylphenyl)-
5 1*H*-pyrazolo[3,4-*b*]pyridin-5-yl]pyrimidin-2-yl]amine;
N-[2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl]-2-
hydroxyacetamide;
N-[2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-
yl]ethyl]piperidine-4-carboxamide;
10 *N*-[2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl]-2-
(methylamino)acetamide;
N-[2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethyl]-2-(2-
hydroxyethylamino)acetamide;
N-[2-[4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-
15 yl]ethyl]nicotinamide;
4-(4-benzyloxyphenyl)-6-(4-fluorophenyl)-2-methyl-5-(4-pyridyl)pyrazolo[3,4-
b]pyridine;
6-(4-fluorophenyl)-5-(2-methylsulfanylpymidin-4-yl)-2-methylpyrazolo[3,4-
b]pyridine;
20 6-(4-fluorophenyl)-2,4-dimethyl-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;
4-(4-benzyloxyphenyl)-6-(4-fluorophenyl)-2-methyl-5-(2-methylsulfanylpymidin-4-
yl)pyrazolo[3,4-*b*]pyridine;
2-(1-benzylpyrrolidin-2-ylmethyl)-4,6-bis(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-
b]pyridine;
25 4-(4-benzyloxyphenyl)-6-(4-fluorophenyl)-2-methyl-5-pyrimidin-4-ylpyrazolo[3,4-
b]pyridine;
6-(4-fluorophenyl)-2-methyl-4-(5-methylfuran-2-yl)-5-(4-pyridyl)pyrazolo[3,4-
b]pyridine;
4-(5-bromofuran-2-yl)-6-(4-fluorophenyl)-2-methyl-5-(4-pyridyl)pyrazolo[3,4-
30 *b*]pyridine;
6-(4-fluorophenyl)-2-methyl-4-propyl-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;
4-(3-benzyloxyphenyl)-6-(4-fluorophenyl)-2-methyl-5-(4-pyridyl)pyrazolo[3,4-
b]pyridine;

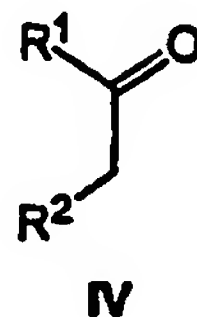
- 6-(4-fluorophenyl)-2-methyl-4-(2-phenylethyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;
5-(2-chloropyridin-4-yl)-6-(4-fluorophenyl)-2-methylpyrazolo[3,4-*b*]pyridine;
4-benzyl-6-(4-fluorophenyl)-2-methyl-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridine;
4-[6-(4-fluorophenyl)-2-methyl-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-4-yl]butan-1-ol;
5 4-[6-(4-fluorophenyl)-2-methyl-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-4-yl]phenol;
N-[6-(4-fluorophenyl)-5-(2-methylsulfanylpirimidin-4-yl)-1*H*-pyrazolo[3,4-*b*]pyridin-3-yl]acetamide;
N-[5-[2-[(cyclopropylmethyl)amino]pyrimidin-4-yl]-6-(4-fluorophenyl)-1*H*-pyrazolo[3,4-*b*]pyridin-3-yl]acetamide;
10 3-[6-(4-fluorophenyl)-4-(2-furyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]propan-1-ol;
2-[4-(4-benzyloxyphenyl)-6-(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]ethanol;
3-[4-(4-benzyloxyphenyl)-6-(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]propan-1-ol;
15 3-[6-(4-fluorophenyl)-4-(5-methylfuran-2-yl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]propan-1-ol;
3-[4-cyclopropyl-6-(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]propan-1-ol;
20 3-[4-(5-bromothien-2-yl)-6-(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]propan-1-ol;
3-[6-(4-fluorophenyl)-4-propyl-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]propan-1-ol;
3-[4,6-bis(4-fluorophenyl)-5-pyrimidin-4-ylpyrazolo[3,4-*b*]pyridin-2-yl]propan-1-ol;
3-[4-(3-benzyloxyphenyl)-6-(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]propan-1-ol;
25 3-[4-benzyl-6-(4-fluorophenyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-2-yl]propan-1-ol;
(1*S*)-*N*-(1-phenylethyl)-[4-[6-(4-fluorophenyl)-3-methyl-1*H*-pyrazolo[3,4-*b*]pyridin-5-yl]pyrimidin-2-yl]amine;
N-cyclopropylmethyl-[4-[6-(4-fluorophenyl)-3-methyl-1*H*-pyrazolo[3,4-*b*]pyridin-5-yl]pyrimidin-2-yl]amine;
30 1-[4-[6-(4-fluorophenyl)-2-methylpyrazolo[3,4-*b*]pyridin-5-yl]pyrimidin-2-ylamino]propan-2-ol;

- N*-cyclopropylmethyl-[4-[6-phenyl-1*H*-pyrazolo[3,4-*b*]pyridin-5-yl]pyrimidin-2-yl]amine;
- 2-[4-[6-(4-fluorophenyl)-2-methylpyrazolo[3,4-*b*]pyridin-5-yl]pyrimidin-2-ylamino]propan-1-ol;
- 5 4-[4-[6-(4-fluorophenyl)-2-methylpyrazolo[3,4-*b*]pyridin-5-yl]pyrimidin-2-ylamino]butan-1-ol;
- (1*S*)-*N*-(1-phenylethyl)-[4-[6-phenyl-1*H*-pyrazolo[3,4-*b*]pyridin-5-yl]pyrimidin-2-yl]amine;
- N*-(3-methoxypropyl)-[4-[6-(3-trifluoromethylphenyl)-1*H*-pyrazolo[3,4-*b*]pyridin-5-yl]pyrimidin-2-yl]amine;
- 10 3-[4-[6-(3-trifluoromethylphenyl)-1*H*-pyrazolo[3,4-*b*]pyridin-5-yl]pyrimidin-2-ylamino]propan-1-ol;
- 3-[4-[6-(4-fluorophenyl)-2-(3-hydroxypropyl)pyrazolo[3,4-*b*]pyridin-5-yl]pyrimidin-2-ylamino]propan-1-ol;
- 15 *N*-ethyl-[4-[6-(3-trifluoromethylphenyl)-1*H*-pyrazolo[3,4-*b*]pyridin-5-yl]pyrimidin-2-yl]amine;
- N*-benzyl-[4-[6-(3-trifluoromethylphenyl)-1*H*-pyrazolo[3,4-*b*]pyridin-5-yl]pyrimidin-2-yl]amine;
- 4-[5-[2-[(cyclopropylmethyl)amino]pyrimidin-4-yl]-6-(4-fluorophenyl)-2-methylpyrazolo[3,4-*b*]pyridin-4-yl]phenol;
- 20 4-[6-(4-fluorophenyl)-2-(3-hydroxypropyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-4-yl]phenol;
- 4-[6-(4-fluorophenyl)-2-methyl-5-pyrimidin-4-ylpyrazolo[3,4-*b*]pyridin-4-yl]phenol;
- 3-[6-(4-fluorophenyl)-2-(3-hydroxypropyl)-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-4-yl]phenol;
- 25 3-[6-(4-fluorophenyl)-2-methyl-5-(4-pyridyl)pyrazolo[3,4-*b*]pyridin-4-yl]phenol;
- 4,6-bis(4-fluorophenyl)-5-(4-pyridyl)-2-(pyrrolidin-2-ylmethyl)pyrazolo[3,4-*b*]pyridine;
- 4-[4-[6-(4-fluorophenyl)-2-methylpyrazolo[3,4-*b*]pyridin-5-yl]pyridin-2-ylamino]benzenesulfonamide;
- 30 *N*-[5-[2-[(cyclopropylmethyl)amino]pyrimidin-4-yl]-6-(4-fluorophenyl)-1*H*-pyrazolo[3,4-*b*]pyridin-3-yl]acetamide 7-oxide; and
- N*-[6-(4-fluorophenyl)-5-(4-pyridyl)-1*H*-pyrazolo[3,4-*b*]pyridin-3-yl]isonicotinamide;

or a salt, solvate or prodrug thereof.

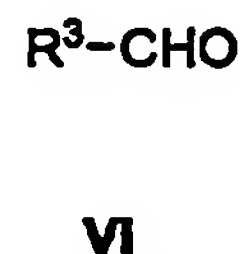
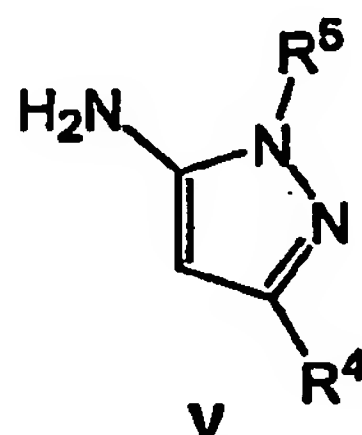
28.- Process for preparing a compound of formula I according to claim 1, which comprises:

(a) reacting a ketone of formula IV



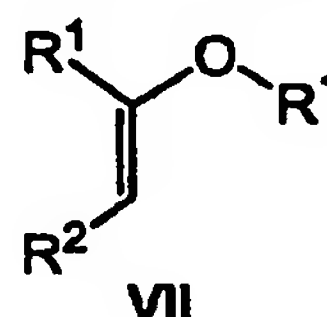
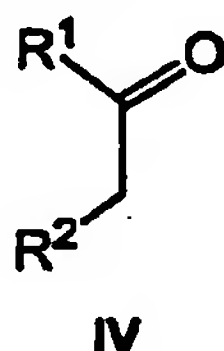
5

wherein R¹ and R² have the same meaning as in claim 1, with an aminopyrazole of formula V and an aldehyde of formula VI

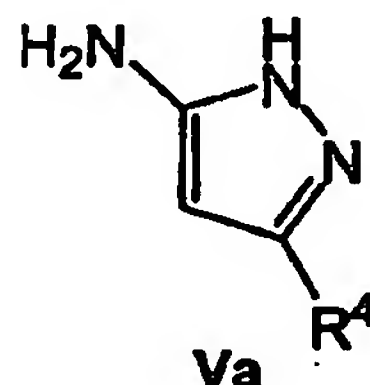


wherein R³, R⁴ and R⁵ have the same meaning as in claim 1; or

10 (b) when in a compound of formula I, R⁵ represents H and R³ has the same meaning as R¹, reacting a ketone of formula IV or an enolate of formula VII



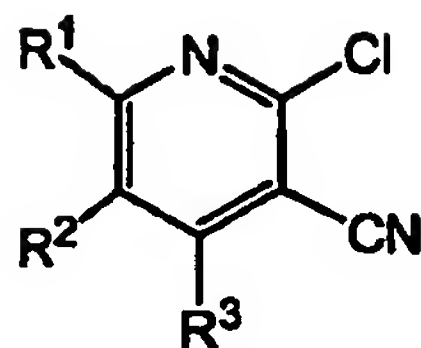
15 wherein R¹ and R² have the same meaning as in claim 1, with an aminopyrazole of formula Va.



wherein R⁴ has the same meaning as in claim 1; or

20 (c) when in a compound of formula I R⁴ represents NH₂, treating a compound of formula XIX

157



XIX

wherein R¹, R² and R³ have the same meaning as in claim 1, with a hydrazine of formula VIIIa



VIIIa

- 5 wherein R⁵ has the same meaning as in claim 1; or
- (d) converting, in one or a plurality of steps, a compound of formula I into another compound of formula I; and
- (e) if desired, after the previous steps, reacting a compound of formula I with a base or an acid to give the corresponding salt.
- 10 29.- A pharmaceutical composition which comprises an effective amount of a compound of formula I according to any of claims 1 to 27 or a pharmaceutically acceptable salt, solvate or prodrug thereof and one or more pharmaceutically acceptable excipients.
- 30.- Use of a compound of formula I according to any of claims 1 to 27 or a pharmaceutically acceptable salt, solvate or prodrug thereof for the manufacture of
- 15 a medicament for the treatment or prevention of diseases mediated by p38.
- 31.- Use of a compound of formula I according to any of claims 1 to 27 or a pharmaceutically acceptable salt, solvate or prodrug thereof for the manufacture of a medicament for the treatment or prevention of diseases mediated by cytokines.
- 20 32.- Use of a compound of formula I according to any of claims 1 to 27 or a pharmaceutically acceptable salt, solvate or prodrug thereof for the manufacture of a medicament for the treatment or prevention of diseases mediated by TNF- α , IL-1, IL-6 and/or IL-8.
- 33.- Use of a compound of formula I according to any of claims 1 to 27 or a pharmaceutically acceptable salt, solvate or prodrug thereof for the manufacture of
- 25 a medicament for the treatment or prevention of a disease selected from immune, autoimmune and inflammatory diseases, cardiovascular diseases, infectious

diseases, bone resorption diseases, neurodegenerative diseases, proliferative diseases and processes associated with the induction of cyclooxygenase-2.

INTERNATIONAL SEARCH REPORT

International Application No
PCT/EP2004/001974

A. CLASSIFICATION OF SUBJECT MATTER
IPC 7 C07D471/04 A61K31/437

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 C07D

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal, CHEM ABS Data, WPI Data, PAJ

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
P,A	WO 03/033502 A (DAVIS JEREMY MARTIN ; BROOKINGS DANIEL CHRISTOPHER (GB); CELLTECH R &) 24 April 2003 (2003-04-24) page 128 - page 134; claims 1-22	1-33
Y	WO 02/062795 A (FUJIMURA MASATAKE ; FUKUNAGA KENJI (JP); HANANO TOKUSHI (JP); OKABE HI) 15 August 2002 (2002-08-15) page 218 - page 224; claims 1-18	1-33
Y	WO 01/53268 A (AGOURON PHARMA) 26 July 2001 (2001-07-26) page 162 - page 182; claims 1-24	1-33
Y	WO 99/30710 A (SQUIBB BRISTOL MYERS CO) 24 June 1999 (1999-06-24) page 30 - page 37; claims 1-42	1-33
	-/-	

☒ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

* Special categories of cited documents:

- *A* document defining the general state of the art which is not considered to be of particular relevance
- *E* earlier document but published on or after the international filing date
- *L* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- *O* document referring to an oral disclosure, use, exhibition or other means
- *P* document published prior to the international filing date but later than the priority date claimed

- *T* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- *X* document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- *Y* document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.
- *8* document member of the same patent family

Date of the actual completion of the international search

6 July 2004

Date of mailing of the international search report

14/07/2004

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,
Fax: (+31-70) 340-3016

Authorized officer

Kyriakakou, G

INTERNATIONAL SEARCH REPORT

International Application No
PCT/EP2004/001974

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT		
Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	US 2002/013328 A1 (SHERK SUSAN R ET AL) 31 January 2002 (2002-01-31) page 43 - page 49; claims 1-8 -----	1-33

INTERNATIONAL SEARCH REPORT

International Application No

PCT/EP2004/001974

Patent document cited in search report		Publication date		Patent family member(s)	Publication date
WO 03033502	A	24-04-2003	CA	2462147 A1	24-04-2003
			WO	03033502 A1	24-04-2003
WO 02062795	A	15-08-2002	CA	2437215 A1	15-08-2002
			EP	1355909 A2	29-10-2003
			WO	02062795 A2	15-08-2002
			US	2004052822 A1	18-03-2004
WO 0153268	A	26-07-2001	AU	2953901 A	31-07-2001
			BG	107011 A	30-04-2003
			BR	0107783 A	19-11-2002
			CA	2388885 A1	26-07-2001
			CN	1394205 T	29-01-2003
			EE	200200398 A	15-10-2003
			EP	1250326 A2	23-10-2002
			HU	0203965 A2	28-05-2003
			JP	2003520273 T	02-07-2003
			NO	20022117 A	16-09-2002
			SK	10052002 A3	04-03-2003
			WO	0153268 A2	26-07-2001
			US	2003139463 A1	24-07-2003
			US	2002161022 A1	31-10-2002
			ZA	200203040 A	11-08-2003
WO 9930710	A	24-06-1999	AU	747705 B2	23-05-2002
			AU	1629799 A	05-07-1999
			CA	2314355 A1	24-06-1999
			EP	1043998 A1	18-10-2000
			JP	2002508324 T	19-03-2002
			WO	9930710 A1	24-06-1999
			US	6107305 A	22-08-2000
			ZA	9811178 A	07-06-2000
US 2002013328	A1	31-01-2002	US	6531477 B1	11-03-2003
			CA	2431038 A1	06-09-2002
			EP	1383769 A2	28-01-2004
			WO	02067654 A2	06-09-2002
			AU	6422399 A	01-05-2000
			CA	2345809 A1	20-04-2000
			EP	1121363 A2	08-08-2001
			JP	2002537223 T	05-11-2002
			WO	0021926 A2	20-04-2000

CORRECTED VERSION

(19) World Intellectual Property
Organization
International Bureau



(43) International Publication Date
10 September 2004 (10.09.2004)

PCT

(10) International Publication Number
WO 2004/076450 A1

(51) International Patent Classification⁷: C07D 471/04,
A61K 31/437

(21) International Application Number:
PCT/EP2004/001974

(22) International Filing Date: 27 February 2004 (27.02.2004)

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data:
ES 2003 00640 27 February 2003 (27.02.2003) ES
ES 2003 00727 14 March 2003 (14.03.2003) ES
ES 2003 02504 27 October 2003 (27.10.2003) ES

(71) Applicant (for all designated States except US): J. URI-
ACH Y COMPAÑIA S.A. [ES/ES]; Av. Camí Reial,
51-57, E-08184 Palau-Solità i Plegamans (ES).

(72) Inventors; and

(75) Inventors/Applicants (for US only): ALMANSA ROS-
ALES, Carmen [ES/ES]; C/Teodora Lamadrid, 41,
E-08022 Barcelona (ES). VIRGILI BERNADÓ, Marina
[ES/ES]; Plaça Lesseps, 11, E-08023 Barcelona (ES).

(74) Agent: ZUMSTEIN F. and KLINGSEISEN F.;
Bräunhausstr. 4, 80331 München (DE).

(81) Designated States (unless otherwise indicated, for every
kind of national protection available): AE, AG, AL, AM,

AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN,
CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI,
GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE,
KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD,
MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG,
PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM,
TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM,
ZW.

(84) Designated States (unless otherwise indicated, for every
kind of regional protection available): ARIPO (BW, GH,
GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW),
Eurasian (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), Euro-
pean (AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR,
GB, GR, HU, IE, IT, LI, MC, NL, PT, RO, SE, SI, SK,
TR), OAPI (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW,
ML, MR, NE, SN, TD, TG).

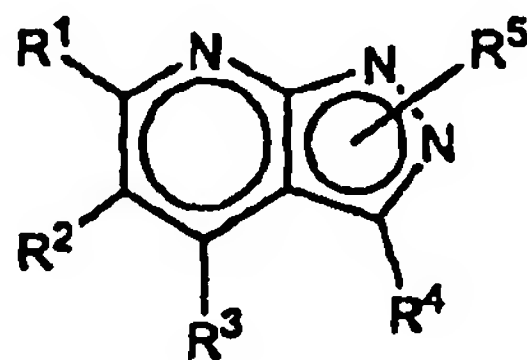
Published:
— with international search report

(48) Date of publication of this corrected version:
1 September 2005

(15) Information about Correction:
see PCT Gazette No. 35/2005 of 1 September 2005, Sec-
tion II

For two-letter codes and other abbreviations, refer to the "Guid-
ance Notes on Codes and Abbreviations" appearing at the begin-
ning of each regular issue of the PCT Gazette.

(54) Title: PYRAZOLOPYRIDINE DERIVATES



(I)

(57) Abstract: New compounds of formula (I) and the salts, solvates and pro-
drugs thereof, wherein the meanings for the various substituents are as disclosed
in the description. These compounds are useful as p38 kinase inhibitors.

WO 2004/076450 A1